=> fil reg FILE 'REGISTRY' ENTERED AT 13:09:25 ON 16 JUL 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 American Chemical Society (ACS)

STRUCTURE FILE UPDATES: 15 JUL 2002 HIGHEST RN 438572-95-3 DICTIONARY FILE UPDATES: 15 JUL 2002 HIGHEST RN 438572-95-3

TSCA INFORMATION NOW CURRENT THROUGH January 7, 2002

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

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L139 ANSWER 1 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **345950-34-7** REGISTRY

CN Uridine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with cytidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H26 N5 O19 P3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

HO

OH

PAGE 1-B

PAGE 1-A

-NH<sub>2</sub>

Jan Delaval
Reference Librarian
siotechnology & Chemical Library
CM1 1E07 – 703-308-4498
jan.delaval@uspto.gov

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

- 3 REFERENCES IN FILE CA (1967 TO DATE)
- 3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:690

REFERENCE 3: 135:71289

L139 ANSWER 2 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN 318250-11-2 REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 2'-deoxycytidine, tetrasodium salt (9CI) (CA INDEX NAME)

OTHER NAMES:

CN INS 37217

FS STEREOSEARCH

MF C18 H27 N5 O21 P4 . 4 Na

SR CA

LC STN Files: CA, CAPLUS, DRUGNL, DRUGUPDATES, USPATFULL

CRN (211448-85-0)

Absolute stereochemistry.

PAGE 1-A

•4 Na

d CR4 U (salt)

PAGE 1-B

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350561

REFERENCE 2: 136:200422

REFERENCE 3: 135:298823

REFERENCE 4: 134:91109

L139 ANSWER 3 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN 211448-85-0 REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 2'-deoxycytidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N5 O21 P4

CI COM

SR CA

LC STN Files: CA, CAPLUS, DRUGUPDATES, TOXCENTER, USPATFULL

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

d CP4U

## \*\*PROPERTY\_DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

9 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

9 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:151393

REFERENCE 3: 136:690

REFERENCE 4: 135:298823

REFERENCE 5: 135:71289

REFERENCE 6: 134:91109

REFERENCE 7: 133:12775

REFERENCE 8: 132:453

REFERENCE 9: 129:175919

L139 ANSWER 4 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **211448-78-1** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), 2'-deoxy-, P'''.fwdarw.5'-ester

with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H26 N4 O22 P4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

PAGE 1-B

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:151393

REFERENCE 3: 136:690

REFERENCE 4: 135:298823

REFERENCE 5: 135:71289

REFERENCE 6: 129:175919

L139 ANSWER 5 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **211448-70-3** REGISTRY

CN Uridine 5'-(hexahydrogen pentaphosphate), P''''.fwdarw.5'-ester with

uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N4 O26 P5

CI COM

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

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PAGE 1-B

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#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:151393

REFERENCE 3: 136:690

REFERENCE 4: 135:298823

REFERENCE 5: 135:71289

REFERENCE 6: 129:175919

L139 ANSWER 6 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **111648-11-4** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with

cytidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N5 O22 P4

CI COM

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

CP4U

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

9 REFERENCES IN FILE CA (1967 TO DATE)

9 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:350561

REFERENCE 3: 136:200422

REFERENCE 4: 136:151393

REFERENCE 5: 136:690

REFERENCE 6: 135:298823

REFERENCE 7: 135:71289

REFERENCE 8: 129:175919

REFERENCE 9: 107:232748

L139 ANSWER 7 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **81534-69-2** REGISTRY

CN Inosine 5'-(hexahydrogen pentaphosphate), P''''.fwdarw.5'-ester with

inosine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H27 N8 O24 P5

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

PAGE 1-A

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PAGE 1-B

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:690

REFERENCE 3: 135:71289

REFERENCE 4: 133:276635

REFERENCE 5: 132:44923

REFERENCE 6: 96:177213

L139 ANSWER 8 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **63785-59-1** REGISTRY

FS STEREOSEARCH

MF C18 H25 N4 O20 P3

CI COM

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

UzP3

8 REFERENCES IN FILE CA (1967 TO DATE)

8 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:151393

REFERENCE 3: 136:690

REFERENCE 4: 135:71289

REFERENCE 5: 132:265443

REFERENCE 6: 130:125350

REFERENCE 7: 129:175919

REFERENCE 8: 88:70843

L139 ANSWER 9 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN 59985-21-6 REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

OTHER NAMES:

CN P1,P4-Diuridine 5'-tetraphosphate

FS STEREOSEARCH

MF C18 H26 N4 O23 P4

CI COM

LC STN Files: CA, CAPLUS, CASREACT, DRUGNL, DRUGPAT, DRUGUPDATES, TOXCENTER, USPATFULL

Absolute stereochemistry.

PAGE 1-B

$$U_2G_4$$

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

20 REFERENCES IN FILE CA (1967 TO DATE)
21 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:151393

REFERENCE 3: 136:690

REFERENCE 4: 135:358114

REFERENCE 5: 135:298823

REFERENCE 6: 135:71289

REFERENCE 7: 134:361372

REFERENCE 8: 133:12775

REFERENCE 9: 132:265443

REFERENCE 10: 131:153747

L139 ANSWER 10 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **30632-06-5** REGISTRY

CN Adenosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-(tetrahydrogen triphosphate), 5'.fwdarw.5'-ester with uridine (8CI)

OTHER NAMES:

CN P1,P3-(Adenosine-5'-uridine-5') triphosphate

FS STEREOSEARCH

DR 83008-72-4

MF C19 H26 N7 O18 P3

LC STN Files: CA, CAPLUS, MEDLINE, TOXCENTER, USPATFULL

Absolute stereochemistry.

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UP3A.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

8 REFERENCES IN FILE CA (1967 TO DATE)

8 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:350567

REFERENCE 2: 136:690

REFERENCE 3: 135:71289

REFERENCE 4: 107:93017

REFERENCE 5: 106:116316

REFERENCE 6: 104:225161

REFERENCE 7: 97:158617

#### REFERENCE 8: 74:19380

L139 ANSWER 11 OF 12 REGISTRY COPYRIGHT 2002 ACS

RN **5959-90-0** REGISTRY

CN Adenosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with adenosine (8CI, 9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-triphosphate, 5'-ester with adenosine (7CI)

OTHER NAMES:

CN Adenosine-(5')-triphospho-(5')-adenosine

CN Ap3A

CN ApppA

CN P1, P3-Diadenosine-5' triphosphate

FS STEREOSEARCH

DR 158700-26-6

MF C20 H27 N10 O16 P3

CI COM

LC STN Files: AGRICOLA, ANABSTR, BEILSTEIN\*, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, TOXCENTER, USPATFULL

(\*File contains numerically searchable property data)

#### Absolute stereochemistry.

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PAGE 1-B

A283

#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

217 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

217 REFERENCES IN FILE CAPLUS (1967 TO DATE)

4 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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1: 137:16915
REFERENCE
REFERENCE
            2:
                136:365598
REFERENCE
                136:350567
            3:
                136:319650
REFERENCE'
            4:
                136:275130
REFERENCE
            5:
REFERENCE
            6:
                136:274624
                136:257619
REFERENCE
            7:
REFERENCE
            8:
                136:151393
                136:144657
REFERENCE
            9:
                136:113039
REFERENCE 10:
L139 ANSWER 12 OF 12 REGISTRY COPYRIGHT 2002 ACS
     5542-28-9 REGISTRY
     Adenosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with
CN
                      (CA INDEX NAME)
     adenosine (9CI)
OTHER CA INDEX NAMES:
     5'-Adenylic acid, 5'.fwdarw.5'-anhydride with adenosine 5'-triphosphate
     Adenosine 5'-(pentahydrogen tetraphosphate), 5'.fwdarw.5'-ester with
CN
     adenosine (8CI)
     Adenosine 5'-tetraphosphate, 5'-ester with adenosine (7CI)
CN
OTHER NAMES:
CN
     5',5'''-Diadenosine tetraphosphate
     Adenosine-5'-tetraphospho-5'-adenosine
CN
CN
     Ap4A
CN
     AppppA
     Diadenosine 5',5'''-P1,P4-tetraphosphate
CN
CN
     Diadenosine tetraphosphate
CN
     P1, P4-Di (adenosin-5'-yl) tetraphosphate
CN
     P1, P4-Diadenosine-5'-tetraphosphate
FS
     STEREOSEARCH
     166104-03-6, 128113-54-2, 138172-62-0, 88109-91-5, 117137-55-0,
DR
     199995-60-3
MF
     C20 H28 N10 O19 P4
CI
     COM
                  AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
LC
       BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB, CIN, EMBASE,
       MEDLINE, RTECS*, TOXCENTER, USPATFULL
         (*File contains numerically searchable property data)
```

Absolute stereochemistry.

AGYA

PAGE 1-B

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

561 REFERENCES IN FILE CA (1967 TO DATE)

18 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

561 REFERENCES IN FILE CAPLUS (1967 TO DATE) 3 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 137:17106

REFERENCE 2: 137:16915

REFERENCE 3: 136:396247

REFERENCE 4: 136:350567

REFERENCE 5: 136:336848

REFERENCE 6: 136:334993

REFERENCE 7: 136:319650

REFERENCE 8: 136:291146

REFERENCE 9: 136:275130

REFERENCE 10: 136:257619

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L141 ANSWER 1 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-23-1 REGISTRY

CN Uridine 5'-(trihydrogen diphosphate), 4-thio-, P'.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H24 N4 O16 P2 S

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

#### Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

L141 ANSWER 2 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-22-0 REGISTRY

CN Uridine 5'-(tetrahydrogen triphosphate), 4-thio-, P''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H25 N4 O19 P3 S

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

#### Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

- 1 REFERENCES IN FILE CA (1967 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

#### REFERENCE 1: 135:298823

L141 ANSWER 3 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-21-9 REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), 6-O-eicosyl-,
P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C39 H66 N6 O22 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

## Absolute stereochemistry.

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

## REFERENCE 1: 135:298823

L141 ANSWER 4 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-20-8 REGISTRY

CN Adenosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 4-(mercaptomethyl)-1-.beta.-D-ribofuranosyl-2(1H)-pyrimidinone (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H29 N7 O20 P4 S

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

L141 ANSWER 5 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-19-5 REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), 4-thio-, P'''.fwdarw.5'-ester with 4-amino-1-.beta.-D-arabinofuranosyl-2(1H)-pyrimidinone (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N5 O21 P4 S

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

L141 ANSWER 6 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-18-4 REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), 6-thio-, P'''.fwdarw.5'-ester with 2'-deoxyuridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 N6 O20 P4 S

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

#### 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

L141 ANSWER 7 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **366004-17-3** REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), 6-thio-, P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 N6 O21 P4 S

CI COM

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

L141 ANSWER 8 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 366004-16-2 REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 3-.beta.-D-ribofuranosyl-3H-1,2,3-triazolo[4,5-d]pyrimidin-7-amine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H26 N8 O21 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

L141 ANSWER 9 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **366004-15-1** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine 2',3'-dibenzoate, 2',3'-dibenzoate (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C46 H42 N4 O27 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

Ph

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 135:298823 REFERENCE

L141 ANSWER 10 OF 81 REGISTRY COPYRIGHT 2002 ACS

251317-44-9 REGISTRY

Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN 2'-deoxycytidine, tetraammonium salt (9CI) (CA INDEX NAME)

FS STEREOSEARCH

C18 H27 N5 O21 P4 . 4 H3 N MF

SR

STN Files: CA, CAPLUS, USPATFULL LC

(211448-85-0) CRN

Absolute stereochemistry.

1 REFERENCES IN FILE CAPILLS (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:453

L141 ANSWER 11 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-88-3** REGISTRY

CN Xanthosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with

xanthosine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H25 N8 O20 P3

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

L141 ANSWER 12 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-84-9** REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), 2'-deoxy-, P'''.fwdarw.5'-ester

with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 N6 O21 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

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PAGE 1-B

#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)
3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 13 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-83-8** REGISTRY

CN Adenosine 5'-(pentahydrogen tetraphosphate), 2'-deoxy-, P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H27 N7 O20 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

# \*\*PROPERTY DATA AVAILABLE IN THE PROP FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 14 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-82-7** REGISTRY

CN Guanosine 5'-(pentahydrogen tetraphosphate), 2'-deoxy-,

P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H27 N7 O21 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

PAGE 1-A

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 15 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-81-6** REGISTRY

CN Uridine 5'-(heptahydrogen hexaphosphate), P''''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H28 N4 O29 P6

CI COM

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

HO

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 16 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-80-5** REGISTRY

CN Thymidine 5'-(pentahydrogen tetraphosphate), 3'-azido-3'-deoxy-,
P''-'.fwdarw.5'-ester with 3'-azido-3'-deoxythymidine (9CI) (CA INDEX-

NAME)

FS STEREOSEARCH

MF C20 H28 N10 O19 P4

SR CF

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

3 REFERENCES IN FILE CA (1967 TO DATE) 3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 136:151393 REFERENCE

REFERENCE 2: 135:298823

REFERENCE 129:175919 3:

L141 ANSWER 17 OF 81 REGISTRY COPYRIGHT 2002 ACS

211448-79-2 REGISTRY

Thymidine 5'-(pentahydrogen tetraphosphate), 3'-azido-3'-deoxy-, P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME) -CN

STEREOSEARCH FS

C19 H27 N7 O21 P4 MF

SR

STN Files: CA, CAPLUS, USPATFULL LC

PAGE 1-B

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 18 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-77-0** REGISTRY

CN Xanthosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 N6 O23 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 19 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-76-9** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 4-amino-1-.beta.-D-arabinofuranosyl-2(1H)-pyrimidinone (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N5 O22 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 20 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-75-8** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), 4-thio-, P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H26 N4 O22 P4 S

CI COM-

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 21 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-74-7** REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 N6 O22 P4

CI- COM

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 22 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-73-6** REGISTRY

CN Thymidine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H28 N4 O22 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 23 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 211448-72-5 REGISTRY

CN Imidazo[1,2-c]pyrimidin-5(6H)-one, 6-[5-O-[hydroxy[[hydroxy[[hydroxy(phosp honooxy)phosphinyl]oxy]phosphinyl]oxy]phosphinyl]-.beta.-D-ribofuranosyl]-2-(3-nitrophenyl)-, P'''.fwdarw.5'-ester with 2-(3-nitrophenyl)-6-.beta.-D-ribofuranosylimidazo[1,2-c]pyrimidin-5(6H)-one, tetraammonium salt (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C34 H34 N8 O25 P4 . 4 H3 N

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

● 4 NH3

PAGE 1-B

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:175919

L141 ANSWER 24 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-71-4** REGISTRY

CN Cytidine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with cytidine, tetraammonium salt (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H28 N6 O21 P4 . 4 H3 N

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

CRN (111035-55-3)

● 4 NH3

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2 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

L141 ANSWER 25 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211448-67-8** REGISTRY

CN 5'-Uridylic acid, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with imidodiphosphoric acid, ammonium salt (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N5 O22 P4 . x H3 N

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

CRN (211427-10-0)

●x NH3

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2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

L141 ANSWER 26 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211427-11-1** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), 4-thio-, P'''.fwdarw.5'-ester with 4-thiouridine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H26 N4 O21 P4 S2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:180137

REFERENCE 4: 129:175919

L141 ANSWER 27 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211427-10-0** REGISTRY

CN 5'-Uridylic acid, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with imidodiphosphoric acid (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H27 N5 O22 P4

CI COM

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

2 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

REFERENCE 2: 129:180137

L141 ANSWER 28 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211427-09-7** REGISTRY

CN 5'-Uridylic acid, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with (difluoromethylene)bis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 F2 N4 O22 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:180137

REFERENCE 4: 129:175919

L141 ANSWER 29 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211427-08-6** REGISTRY

CN Uridine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine, tetrasodium salt (9CI) (CA INDEX NAME)

OTHER NAMES:

CN INS 365

CN P1, P4-Diuridine 5'-tetraphosphate tetrasodium salt

FS STEREOSEARCH

DR 266356-23-4

MF C18 H26 N4 O23 P4 . 4 Na

SR CA

LC STN Files: BIOSIS, CA, CAPLUS, CASREACT, DRUGPAT, DRUGUPDATES, PHAR, TOXCENTER, USPATFULL

CRN (59985-21-6)

●4 Na

PAGE 1-B

12 REFERENCES IN FILE CA (1967 TO DATE)
13 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:200422

REFERENCE 2: 136:164777

REFERENCE 3: 136:47750

REFERENCE 4: 136:705

REFERENCE 5: 135:358114

REFERENCE 6: 135:142283

REFERENCE 7: 135:87137

REFERENCE 8: 134:252575

REFERENCE 9: 132:317843

REFERENCE 10: 132:265443

L141 ANSWER 30 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211427-07-5** REGISTRY

CN Imidazo[1,2-c]pyrimidin-5(6H)-one, 6-[5-0-[hydroxy[[hydroxy[[hydroxy(phosp

honooxy)phosphinyl]oxy]phosphinyl]-.beta.-D-ribofuranosyl]-, P'''.fwdarw.5'-ester with 6-.beta.-D-ribofuranosylimidazo[1,2-c]pyrimidin-5(6H)-one (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C22 H28 N6 O21 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

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PAGE 1-B

#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:180137

REFERENCE 4: 129:175919

L141 ANSWER 31 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **211427-06-4** · REGISTRY

CN 5'-Uridylic acid, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with methylenebis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H28 N4 O22 P4

SR CA

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:298823

REFERENCE 3: 129:180137

REFERENCE 4: 129:175919

L141 ANSWER 32 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 188560-02-3 REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with

inosine (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Diinosine tetraphosphate

FS STEREOSEARCH

MF C20 H26 N8 O21 P4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:690

REFERENCE 2: 135:298493

REFERENCE 3: 132:44923

REFERENCE 4: 126:233761

L141 ANSWER 33 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 170638-62-7 REGISTRY

CN Guanosine 5'-(pentahydrogen tetraphosphate), N,N,7-trimethyl-, P'''.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 2-(dimethylamino)-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-99-6

MF C23 H35 N10 O21 P4

SR CA

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\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 123:340747

L141 ANSWER 34 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **170638-61-6** REGISTRY

OTHER CA INDEX NAMES:

CN Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 6,9-dihydro-7-methyl-2-(methylamino)-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-98-5

MF C22 H33 N10 O21 P4

SR CA

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HO

\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 123:340747

L141 ANSWER 35 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **170638-60-5** REGISTRY

CN Guanosine 5'-(tetrahydrogen triphosphate), N,N,7-trimethyl-, P''.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Guanosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with 2-(dimethylamino)-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-90-7

MF C23 H34 N10 O18 P3

SR CA

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## \*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 127:343026

REFERENCE 4: 123:340747

L141 ANSWER 36 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 170638-59-2 -REGISTRY

CN Guanosine 5'-(pentahydrogen tetraphosphate), 7-methyl-,

P'''.fwdarw.5'-ester with 7-methylguanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Purinium, 2-amino-6,9-dihydro-9-[5-0-[hydroxy[[hydroxy[[hydroxy(phospho nooxy)phosphinyl]oxy]phosphinyl]oxy]phosphinyl]-.beta.-D-ribofuranosyl]-7-methyl-6-oxo-, inner salt, P'''.fwdarw.5'-ester with 2-amino-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-92-9

MF C22 H34 N10 O21 P4

SR CA

LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

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\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 123:340747

L141 ANSWER 37 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 170638-58-1 REGISTRY

CN Guanosine 5'-(pentahydrogen tetraphosphate), 7-methyl-,

P'''.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with 2-amino-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-97-4

MF C21 H31 N10 O21 P4

SR CA

LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

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\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 127:343026

REFERENCE 4: 123:340747

L141 ANSWER 38 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **170638-57-0** REGISTRY

CN Guanosine 5'-(trihydrogen diphosphate), 7-methyl-, P'.fwdarw.5'-ester with 7-methylguanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Purinium, 2-amino-6,9-dihydro-9-[5-O-[hydroxy(phosphonooxy)phosphinyl]-.beta.-D-ribofuranosyl]-7-methyl-6-oxo-, P'.fwdarw.5'-ester with 2-amino-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-86-1

MF C22 H32 N10 O15 P2

CI COM

SR . CA

LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

Absolute stereochemistry.

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\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 123:340747

L141 ANSWER 39 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **170638-56-9** REGISTRY

CN Guanosine 5'-(trihydrogen diphosphate), 7-methyl-, P'.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Guanosine 5'-(trihydrogen diphosphate), P'.fwdarw.5'-ester with 2-amino-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-87-2

MF C21 H29 N10 O15 P2

CI COM

SR CA

LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

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### \*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 127:343026

REFERENCE 4: 123:340747

L141 ANSWER 40 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **154960-70-0** REGISTRY

CN Guanosine 5'-(tetrahydrogen triphosphate), 7-methyl-, P''.fwdarw.5'-ester with 7-methylguanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Purinium, 2-amino-6,9-dihydro-7-methyl-6-oxo-9-.beta.-D-ribofuranosyl-, 5'.fwdarw.P:5'''.fwdarw.P''-(trihydrogen triphosphate)

FS STEREOSEARCH

DR 211448-89-4

MF C22 H33 N10 O18 P3

SR - CA

LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

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\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 123:340747

REFERENCE 4: 120:299198

L141 ANSWER 41 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 148503-84-8 REGISTRY

CN Inosine 5'-(pentahydrogen tetraphosphate), P.L. fwdarw.5'-ester with adenosine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H27 N9 O20 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

7 REFERENCES IN FILE CA (1967 TO DATE)
7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:298823

REFERENCE 2: 129:287192

REFERENCE 3: 127:95515

REFERENCE 4: 125:215341

REFERENCE 5: 125:136013

REFERENCE 6: 120:8922

REFERENCE 7: 119:49812

L141 ANSWER 42 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 135802-64-1 REGISTRY

CN 5'-Adenylic acid, 2-amino-, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with (dichloromethylene)bis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H30 C12 N12 O18 P4

SR CA

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OH

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 115:114980

L141 ANSWER 43 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 135780-92-6 REGISTRY

CN 5'-Adenylic acid, 2-amino-, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with methylenebis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H32 N12 O18 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

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### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 115:114980

L141 ANSWER 44 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 135780-85-7 REGISTRY

CN 5'-Adenylic acid, 2-amino-, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with (difluoromethylene)bis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H30 F2 N12 O18 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

OH

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 115:114980

L141 ANSWER 45 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **135780-83-5** REGISTRY

CN Adenosine 5'-(pentahydrogen tetraphosphate), 2-amino-,

P'''.fwdarw.5'-ester with 2-aminoadenosine (9CI) -- (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H30 N12 O19 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 118:250254

REFERENCE 4: 115:114980

L141 ANSWER 46 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **134311-47-0** REGISTRY

CN Guanosine 5'-(tetrahydrogen triphosphate), N,7-dimethyl-, P''.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Guanosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with 6,9-dihydro-7-methyl-2-(methylamino)-6-oxo-9-.beta.-D-ribofuranosyl-1H-purinium

FS STEREOSEARCH

DR 211448-91-8

MF C22 H32 N10 O18 P3

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

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\*\*\* FRAGMENT DIAGRAM IS INCOMPLETE \*\*\*

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 123:340747

REFERENCE 4: 115:44346

L141 ANSWER 47 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **111035-55-3** REGISTRY

CN Cytidine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with cytidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H28 N6 O21 P4

CI COM

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-B

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1967 TO DATE)
3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 107:232748

L141 ANSWER 48 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **103137-89-9** REGISTRY

CN Adenosine 5'-(heptahydrogen hexaphosphate), P''''.fwdarw.5'-ester with thymidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H31 N7 O26 P6

CI COM

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

6 REFERENCES IN FILE CA (1967 TO DATE) 6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 119:176516

REFERENCE 4: 108:182674

REFERENCE 5: 105:221543

REFERENCE 6: 105:38068

L141 ANSWER 49 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **103137-88-8** REGISTRY

CN Adenosine 5'-(hexahydrogen pentaphosphate), P'''.fwdarw.5'-ester with thymidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

DR 204640-72-2

MF C20 H30 N7 O23 P5

CI COM

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

- 9 REFERENCES IN FILE CA (1967 TO DATE)
- 3 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
- 9 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 133:234384

REFERENCE 3: 130:121340

REFERENCE 4: 129:175919

REFERENCE 5: 128:227823

REFERENCE 6: 119:176516

REFERENCE 7: 108:182674

REFERENCE 8: 105:221543

REFERENCE 9: 105:38068

L141 ANSWER 50 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **97776-55-1** REGISTRY

CN 5'-Adenylic acid, 5'.fwdarw.P:5'''.fwdarw.P'-dianhydride with (difluoromethylene)bis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H28 F2 N10 O18 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

9 REFERENCES IN FILE CA (1967 TO DATE)

9 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393-

REFERENCE 2: 129:175919

REFERENCE 3: 118:250254

REFERENCE 4: 113:54862

REFERENCE 5: 112:70015

REFERENCE 6: 111:190199

REFERENCE 7: 110:188318

REFERENCE 8: 109:124879

REFERENCE 9: 103:88157

L141 ANSWER 51 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **97776-54-0** REGISTRY

CN 5'-Adenylic acid, 5'.fwdarw.P,5'''.fwdarw.P'-dianhydride with (dichloromethylene)bis[phosphonic acid] (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H28 C12 N10 O18 P4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP'-FORMAT\*\*

10 REFERENCES IN FILE CA (1967 TO DATE)

10 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 118:250254

REFERENCE 4: 113:54862

REFERENCE 5: 112:70015

REFERENCE 6: 111:190199

REFERENCE 7: 110:188318

REFERENCE 8: 109:124879

REFERENCE 9: 108:127515

REFERENCE 10: 103:88157

L141 ANSWER 52 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **96920-51-3** REGISTRY

CN 3H-Imidazo[2,1-i]purine, 3-[5-O-[hydroxy[[hydroxy[[hydroxy(phosphonooxy)phosphinyl]oxy]phosphinyl]-.beta.-D-ribofuranosyl]-,
P'''.fwdarw.5'-ester with 3-.beta.-D-ribofuranosyl-3H-imidazo[2,1-i]purine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

DR 143721-41-9

MF C24 H28 N10 O19 P4

LC STN Files: CA, CAPLUS, MEDLINE, TOXCENTER, USPATFULL

### Absolute stereochemistry.

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HO

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\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

15 REFERENCES IN FILE CA (1967 TO DATE)

15 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 135:339673

REFERENCE 3: 135:298823

REFERENCE 4: 129:175919

REFERENCE 5: 127:344236

REFERENCE 6: 124:80241

REFERENCE 7: 123:105909

REFERENCE 8: 119:49812

REFERENCE 9: 117:207601

REFERENCE 10: 117:22607

L141 ANSWER 53 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **88109-92-6** REGISTRY

CN 5'-Adenylic acid, 5'.fwdarw.P,5'''.fwdarw.P'-dianhydride with methylenebis[phosphonic acid] (9CI) (CA INDEX NAME)

OTHER NAMES:

CN AppCH2ppA

FS STEREOSEARCH

MF C21 H30 N10 O18 P4

CI COM

LC STN Files: CA, CAPLUS, CASREACT, TOXCENTER, USPATFULL

#### Absolute stereochemistry.

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#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

28 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

28 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 137:17106 REFERENCE

REFERENCE 2: 136:382129

REFERENCE 3: 136:151393

133:134263 REFERENCE 4:

REFERENCE 5: 130:52684

129:175919 REFERENCE 6:

127:95515 REFERENCE 7:

REFERENCE 8: 125:215341

REFERENCE 9: 125:136013

REFERENCE 10: 119:46107

L141 ANSWER 54 OF 81 REGISTRY COPYRIGHT 2002 ACS

83008-69-9 REGISTRY

Adenosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN

cytidine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

148503-81-5, 100849-66-9 DR

C19 H28 N8 O20 P4 MF

CA, CAPLUS, MEDLINE, USPATFULL STN Files:

# Absolute stereochemistry.

#### PAGE 1-A

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### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

19 REFERENCES IN FILE CA (1967 TO DATE)

19 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:365598

REFERENCE 2: 136:151393

135:298823 REFERENCE 3:

REFERENCE 4: 132:264218

REFERENCE 5: 129:287192

REFERENCE 129:175919 6:

REFERENCE 7: 129:132936

125:215341 REFERENCE 8:

120:8922 REFERENCE 9:

REFERENCE 10: 119:49812

L141 ANSWER 55 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **79695-25-3** REGISTRY

Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN (CA INDEX NAME)

uridine (9CI)

STEREOSEARCH FS

C19 H27 N7 O22 P4 MF

STN Files: CA, CAPLUS, USPATFULL LC

Absolute stereochemistry.

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#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

8 REFERENCES IN FILE CA (1967 TO DATE)

8 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 136:151393 REFERENCE

REFERENCE 2: 129:175919

REFERENCE 3: 114:222478

REFERENCE 4: 107:232748

107:54628 REFERENCE 5:

101:19603 REFERENCE 6:

96:47681 REFERENCE 7:

95:182545 REFERENCE 8:

L141 ANSWER 56 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **79695-24-2** REGISTRY

Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN (CA INDEX NAME)

cytidine (9CI)

STEREOSEARCH FS

C19 H28 N8 O21 P4 MF

CA, CAPLUS, USPATFULL LC STN Files:

Absolute stereochemistry.

PAGE 1-A

### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

7 REFERENCES IN FILE CA (1967 TO DATE)
7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 107:232748

REFERENCE 4: 107:54628

REFERENCE 5: 101:19603

REFERENCE 6: 96:47681

REFERENCE 7: 95:182545

L141 ANSWER 57 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **61340-12-3** REGISTRY

CN Guanosine 5'-(tetrahydrogen triphosphate), 7,8-dihydro-7-methyl-, P''.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H31 N10 O18 P3

LC STN Files: CA, CAPLUS, CASREACT, USPATFULL

Absolute stereochemistry.

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1967 TO DATE)

5 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 136:151393 REFERENCE

134:37905 REFERENCE 2:

REFERENCE 133:318884

REFERENCE 129:175919

86:43958 REFERENCE 5:

L141 ANSWER 58 OF 81 REGISTRY COPYRIGHT 2002 ACS

**59985-20-5** REGISTRY

Xanthosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN

xanthosine (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H26 N8 O23 P4

CA, CAPLUS, USPATFULL LC STN Files:

Absolute stereochemistry.

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#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1967 TO DATE) 5 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 131:70274

REFERENCE 3: 129:175919

REFERENCE 4: 97:122648

REFERENCE 5: 85:42852

L141 ANSWER 59 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **56983-23-4** REGISTRY

CN Adenosine 5'-(heptahydrogen hexaphosphate), P''''.fwdarw.5'-ester with adenosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-hexaphosphate, 5'-ester with adenosine (7CI)

OTHER NAMES:

CN Adenosine-(5')-hexaphospho-(5')-adenosine

CN AppppppA

CN Diadenosine hexaphosphate

FS STEREOSEARCH

DR 123396-55-4, 136105-87-8, 199995-64-7

MF C20 H30 N10 O25 P6

CI COM

LC STN Files: AGRICOLA, BEILSTEIN\*, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, EMBASE, MEDLINE, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

117 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

117 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:319650

REFERENCE 2: 136:291146

REFERENCE 3: 136:257619

REFERENCE 4: -136:151393

REFERENCE 5: 136:144657

REFERENCE 6: 136:113039

REFERENCE 7: 136:968

REFERENCE 8: 135:236901

REFERENCE 9: 135:209315

REFERENCE 10: 135:192374

```
L141 ANSWER 60 OF 81 REGISTRY COPYRIGHT 2002 ACS
     41708-91-2 REGISTRY
RN
     Adenosine 5'-(hexahydrogen pentaphosphate), P''''.fwdarw.5'-ester with
CN
     adenosine (9CI)
                      (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Adenosine 5'-pentaphosphate, 5'-5'-ester with adenosine (7CI)
CN
OTHER NAMES:
     Adenosine-(5')-pentaphospho-(5')-adenosine
CN
CN
     ApppppA
CN
     Di(adenosine-5')pentaphosphate
     Diadenosine pentaphosphate
CN
     P1, P5-Bis-(5'-adenosyl) pentaphosphate
CN
CN
     P1, P5-Di (adenosine-5') pentaphosphate
     P1, P5-Diadenosine pentaphosphate
CN
FS
     STEREOSEARCH
     128113-55-3, 117137-56-1, 199995-62-5
DR
ΜF
     C20 H29 N10 O22 P5
CI
     COM
                  AGRICOLA, BEILSTEIN*, BIOSIS, BIOTECHNO, CA, CAOLD, CAPLUS,
LC
     STN Files:
       CASREACT, CHEMCATS, CHEMLIST, CSCHEM, EMBASE, TOXCENTER, USPATFULL
         (*File contains numerically searchable property data)
                      EINECS**
     Other Sources:
         (**Enter CHEMLIST File for up-to-date regulatory information)
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#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

292 REFERENCES IN FILE CA (1967 TO DATE)

14 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

293 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:319650

REFERENCE 2: 136:291146

REFERENCE 3: 136:275130

REFERENCE 4: 136:257619

REFERENCE 5: 136:227153

REFERENCE 6: 136:151393

REFERENCE 7: 136:144657

REFERENCE 8: 136:114723

REFERENCE 9: 136:113039

REFERENCE 10: 136:112435

L141 ANSWER 61 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **34983-48-7** REGISTRY

CN Adenosine 5'-(trihydrogen diphosphate), 2-(methylthio)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-(Methylthio)adenosine 5'-diphosphate

CN 2-Methylthio-ADP

FS STEREOSEARCH

DR 41036-37-7

MF C11 H17 N5 O10 P2 S

LC STN Files: BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, MEDLINE, TOXCENTER, USPATFULL

Absolute stereochemistry.

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

77 REFERENCES IN FILE CA (1967 TO DATE)

77 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 137:28390

REFERENCE 2: 136:380497

REFERENCE 3: 136:363807

REFERENCE 4: 136:350567

REFERENCE 5: 136:304342

REFERENCE 6: 136:129201

REFERENCE 7: 136:112610

REFERENCE 8: 136:48564

REFERENCE 9: 136:15546

REFERENCE 10: 136:15521

L141 ANSWER 62 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **34692-44-9** REGISTRY

CN Guanosine 5'-(trihydrogen diphosphate), P'.fwdarw.5'-ester with guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Guanosine 5'-(trihydrogen pyrophosphate), 5'.fwdarw.5'-ester with guanosine (8CI)

OTHER NAMES:

CN Diguanosine 5',5'''-diphosphate

FS STEREOSEARCH

MF C20 H26 N10 O15 P2

CI COM

LC STN Files: BEILSTEIN\*, CA, CAPLUS, CASREACT, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry.

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## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

- 29 REFERENCES IN FILE CA (1967 TO DATE)
  - 1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
- 29 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 136:144657

REFERENCE 3: 136:65956

REFERENCE 4: 135:253549

REFERENCE 5: 135:3938

REFERENCE 6: 132:50203

REFERENCE 7: 129:175919

REFERENCE 8: 129:149163

REFERENCE 9: 127:190963

REFERENCE 10: 125:196204

L141 ANSWER 63 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **30632-08-7** REGISTRY

CN Adenosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with thymidine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-(tetrahydrogen triphosphate), 5'.fwdarw.5'-ester with thymidine (8CI)

OTHER NAMES:

CN ApppdT

CN P1, P3-(Adenosine-5'-deoxythymidine-5') triphosphate

FS STEREOSEARCH

DR 83028-14-2

MF C20 H28 N7 O17 P3

CI COM

LC STN Files: BEILSTEIN\*, CA, CAPLUS, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry.

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## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

8 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

8 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393

REFERENCE 2: 129:175919

REFERENCE 3: 109:149968

REFERENCE 4: 109:6867

REFERENCE 5: 108:182674

REFERENCE 6: 105:38068

REFERENCE 7: 97:158617

REFERENCE 8: 74:19380

L141 ANSWER 64 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **27821-45-0** REGISTRY

CN Uridine 5'-(trihydrogen diphosphate), disodium salt (8CI, 9CI) (CA INDEX NAME)

OTHER NAMES:

CN UDP disodium salt

CN Uridine-5'-diphosphate disodium salt

FS STEREOSEARCH

MF C9 H14 N2 O12 P2 . 2 Na

LC STN Files: CA, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CSCHEM, IFICDB, IFIPAT, IFIUDB, MSDS-OHS, TOXCENTER, USPATFULL

Other Sources: EINECS\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

CRN (58-98-0)

#### ●2 Na

18 REFERENCES IN FILE CA (1967 TO DATE)
19 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:705

REFERENCE 2: 135:358114

REFERENCE 3: 134:252575

REFERENCE 4: 130:125350

REFERENCE 5: 117:70247

REFERENCE 6: 106:214302

REFERENCE 7: 89:152782

REFERENCE 8: 88:101148

REFERENCE 9: 88:55097

REFERENCE 10: 87:44270

L141 ANSWER 65 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **26184-65-6** REGISTRY

CN Uridine 5'-(trihydrogen diphosphate), P'.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Uridine 5'-(trihydrogen pyrophosphate), 5'.fwdarw.5'-ester with uridine (8CI)

CN Uridine 5'-pyrophosphate, 5'.fwdarw.5'-ester with uridine (7CI)

. . . . . .

CN Uridine pyrophosphate, 5',5'-ester with uridine (6CI)

OTHER NAMES:

CN Uridine diphosphouridine

FS STEREOSEARCH

MF C18 H24 N4 O17 P2

CI COM

LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS, CASREACT, USPATFULL (\*File contains numerically searchable property data)

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

13 REFERENCES IN FILE CA (1967 TO DATE)
13 REFERENCES IN FILE CAPLUS (1967 TO DATE)
6 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:151393

REFERENCE 2: 132:265443

REFERENCE 3: 132:251360

REFERENCE 4: 129:175919

REFERENCE 5: 129:149163

REFERENCE 6: 127:190963

REFERENCE 7: 125:241071

REFERENCE 8: 117:70234

REFERENCE 9: 116:59878

REFERENCE 10: 97:35034

L141 ANSWER 66 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **19817-92-6** REGISTRY

CN Uridine 5'-(tetrahydrogen triphosphate), trisodium salt (8CI, 9CI) (CA INDEX NAME)

OTHER NAMES:

CN Trisodium UTP

CN Uridine-5'-triphosphate trisodium salt

CN Utipina

CN UTP trisodium salt

FS STEREOSEARCH

MF C9 H15 N2 O15 P3 . 3 Na

LC STN Files: BEILSTEIN\*, CA, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CSCHEM, MSDS-OHS, TOXCENTER, USPATFULL

(\*File contains numerically searchable property data)

Other Sources: EINECS\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

CRN (63-39-8)

#### ●3 Na

CRN

(58-64-0)

# 21 REFERENCES IN FILE CA (1967 TO DATE) 22 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:151393 REFERENCE 2: 136:705 135:358114 REFERENCE 3: 134:252575 REFERENCE REFERENCE 5: 130:125350 130:115035 REFERENCE 6: 129:175919 REFERENCE 7: REFERENCE 8: 127:31151 REFERENCE 9: 126:229645 REFERENCE 10: 117:157673 L141 ANSWER 67 OF 81 REGISTRY COPYRIGHT 2002 ACS **16178-48-6** REGISTRY Adenosine 5'-(trihydrogen diphosphate), disodium salt (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: Adenosine-5'-(trihydrogen pyrophosphate), disodium salt (8CI) OTHER NAMES: Adenosine-5'-diphosphate disodium salt CN ADP disodium CN ADP disodium salt CN CN Disodium 5'-ADP Disodium adenosine 5'-diphosphate CN Disodium ADP CN CN Trinosin S FS STEREOSEARCH DR 2921-83-7 C10 H15 N5 O10 P2 . 2 Na MF AGRICOLA, BEILSTEIN\*, BIOBUSINESS, BIOSIS, CA, CAPLUS, LC CASREACT, CHEMCATS, CHEMLIST, CSCHEM, MSDS-OHS, TOXCENTER, USPATFULL (\*File contains numerically searchable property data) EINECS\*\*, NDSL\*\*, TSCA\*\* Other Sources: (\*\*Enter CHEMLIST File for up-to-date regulatory information)

Absolute stereochemistry.

# •2 Na

43 REFERENCES IN FILE CA (1967 TO DATE)

43 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:705

REFERENCE 2: 133:175213

REFERENCE 3: 132:177247

REFERENCE 4: 132:162719

REFERENCE 5: 123:75462

REFERENCE 6: 120:289156

REFERENCE 7: 119:85054

REFERENCE 8: 118:75514

REFERENCE 9: 117:60392

REFERENCE 10: 117:19587

L141 ANSWER 68 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **14264-46-1** REGISTRY

CN Uridine 5'-(tetrahydrogen triphosphate), tetrasodium salt (8CI, 9CI) (CA INDEX NAME)

OTHER NAMES:

CN Sodium UTP

CN Uridine triphosphate sodium salt

FS STEREOSEARCH

MF  $\,$  C9 H15 N2 O15 P3 . 4 Na

LC STN Files: BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, USPATFULL (\*File contains numerically searchable property data)

CRN (63-39-8)

#### Na

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:339428

REFERENCE 2: 86:66745

REFERENCE 3: 86:21792

L141 ANSWER 69 OF 81 REGISTRY COPYRIGHT 2002 ACS

13457-68-6 REGISTRY

Adenosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN

thymidine (8CI, 9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

Adenosine 5'-tetraphosphate, 5'.fwdarw.5'-ester with thymidine (7CI) OTHER NAMES:

CN P1, P4-(Adenosine-5'-deoxythymidine-5') tetraphosphate

Thymidine, 5'-ester with adenosine 5'-tetraphosphate CN

FS STEREOSEARCH

148503-83-7 DR

C20 H29 N7 O20 P4 MF

CI COM

LCSTN Files: CA, CANCERLIT, CAOLD, CAPLUS, MEDLINE, TOXCENTER, USPATFULL

Absolute stereochemistry.

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

14 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

14 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:365598

REFERENCE 2: 136:151393

REFERENCE 3: 132:321022

REFERENCE 4: 129:175919

REFERENCE 5: 129:132936

REFERENCE 6: 125:215341

REFERENCE 7: 120:8922

REFERENCE 8: 119:176516

REFERENCE 9: 119:49812

REFERENCE 10: 108:182674

L141 ANSWER 70 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 10527-48-7 REGISTRY

CN Adenosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with uridine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-(pentahydrogen tetraphosphate), 5'.fwdarw.5'-ester with uridine (8CI)

CN Adenosine tetraphosphate, 5'.fwdarw.5'-ester with uridine (7CI)

CN Uridine, 5'-ester with adenosine 5'-tetraphosphate (8CI)

OTHER NAMES:

CN P1, P4-(Adenosine-5'-uridine-5') tetraphosphate

FS STEREOSEARCH

DR 148503-80-4, 83008-70-2

MF C19 H27 N7 O21 P4

LC STN Files: CA, CANCERLIT, CAOLD, CAPLUS, MEDLINE, USPATFULL

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## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

22 REFERENCES IN FILE CA (1967 TO DATE) 22 REFERENCES IN FILE CAPLUS (1967 TO DATE) 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 136:365598 1:

REFERENCE 136:151393

REFERENCE 135:298823

REFERENCE 132:264217

REFERENCE 129:287192

REFERENCE 129:175919

REFERENCE 129:132936

REFERENCE 125:215341 8:

120:239424 REFERENCE 9:

REFERENCE 10: 120:8922

L141 ANSWER 71 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **10527-46-5** REGISTRY

Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN (CA INDEX NAME) adenosine (9CI)

OTHER CA INDEX NAMES:

Guanosine 5'-(pentahydrogen tetraphosphate), 5'.fwdarw.5'-ester with adenosine (8CI)

Guanosine 5'-tetraphosphate, 5'-ester with adenosine (7CI)

OTHER NAMES:

CN P1,P4-(Adenosine-5'-guanosine-5') tetraphosphate

CN P1-(Adenosine-5' P4-guanosine-5') tetraphosphate

FS STEREOSEARCH

DR 148503-78-0

MF C20 H28 N10 O20 P4

CI COM

LC STN Files: CA, CAOLD, CAPLUS, CASREACT, TOXCENTER, USPATFULL

# Absolute stereochemistry.

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

53 REFERENCES IN FILE CA (1967 TO DATE)

53 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO-1967)

REFERENCE 1: 136:365598

REFERENCE 2: 136:151393

REFERENCE 3: 136:144657

REFERENCE 4: 134:188297

REFERENCE 5: 133:27916

REFERENCE 6: 132:246125

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REFERENCE
                 132:179669
             7:
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REFERENCE 8: 132:179662

REFERENCE 9: 131:256400

REFERENCE 10: 131:70274

L141 ANSWER 72 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **6674-45-9** REGISTRY

Guanosine 5'-(tetrahydrogen triphosphate), P''.fwdarw.5'-ester with CN guanosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

Guanosine 5'-(tetrahydrogen triphosphate), 5'.fwdarw.5'-ester with CN

quanosine (8CI)

Guanosine triphosphate, 5'.fwdarw.5'-ester with guanosine (7CI) CN

Guanosine, 5'-ester with guanosine 5'-(tetrahydrogen triphosphate) (8CI) CN OTHER NAMES:

Diguanosine 5',5'''-triphosphate CN

Diguanosine 5'-triphosphate CN

CN Diguanosine triphosphate

CN Gp3G

CN P1, P3-Diguanosine 5'-triphosphate

STEREOSEARCH FS

79192-45-3 DR

MF C20 H27 N10 O18 P3

CI COM

BEILSTEIN\*, BIOSIS, CA, CAOLD, CAPLUS, CASREACT, CSCHEM, LCSTN Files: IFICDB, IFIPAT, IFIUDB, MEDLINE, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry.

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# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

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67 REFERENCES IN FILE CA (1967 TO DATE)
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3 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

67 REFERENCES IN FILE CAPLUS (1967 TO DATE)

3 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:151393

REFERENCE 136:144657

REFERENCE 3: 135:269163

135:253549 REFERENCE

REFERENCE 5: 135:73117

REFERENCE 6: 134:350916

REFERENCE 7: 132:218454

8: REFERENCE 132:75952

131:164974 REFERENCE 9:

REFERENCE 10: 131:97001

L141 ANSWER 73 OF 81 REGISTRY COPYRIGHT 2002 ACS

4130-19-2 REGISTRY RN

Guanosine 5'-(pentahydrogen tetraphosphate), P'''.fwdarw.5'-ester with CN (CA INDEX NAME) guanosine (9CI)

OTHER CA INDEX NAMES:

Guanosine 5'-(pentahydrogen tetraphosphate), 5'.fwdarw.5'-ester with CN quanosine (8CI)

Guanosine 5'-tetraphosphate, 5'-ester with guanosine (7CI) CN

OTHER NAMES:

Diguanosine 5',5'''-tetraphosphate CN

Diguanosine 5'-tetraphosphate CN

P1, P4-Diguanosine 5'-tetraphosphate CN

FS STEREOSEARCH

DR 79202-52-1

MF C20 H28 N10 O21 P4

CI COM

BEILSTEIN\*, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, EMBASE, LC STN Files: MEDLINE, TOXCENTER, USPATFULL

(\*File contains numerically searchable property data)

Absolute stereochemistry.

# PAGE 1-B

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

82 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

82 REFERENCES IN FILE CAPLUS (1967 TO DATE)

5 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:151393

REFERENCE 2: 136:144657

REFERENCE 3: 135:253549

REFERENCE 4: 134:350916

REFERENCE 5: 134:188297

REFERENCE 6: 133:155161

REFERENCE 7: 133:27916

REFERENCE 8: 131:164974

REFERENCE 9: 131:97001

REFERENCE 10: 130:278412

L141 ANSWER 74 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **2596-55-6** REGISTRY

CN Adenosine 5'-(trihydrogen diphosphate), P'.fwdarw.5'-ester with adenosine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-(trihydrogen pyrophosphate), 5'.fwdarw.5'-ester with adenosine (8CI)

CN Adenosine 5'-diphosphate, 5'.fwdarw.5'-ester with adenosine (6CI)

CN Adenosine 5'-pyrophosphate, 5'-ester with adenosine (7CI)

OTHER NAMES:

CN Diadenosine 5',5'-pyrophosphate

CN Diadenosine 5'-pyrophosphate

CN Diadenosine pyrophosphate

CN P1, P2-Di (adenosine-5') diphosphate

FS STEREOSEARCH

DR 63266-82-0, 221169-23-9, 381211-20-7

MF C20 H26 N10 O13 P2

CI COM

LC STN Files: BEILSTEIN\*, BIOSIS, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, MEDLINE, TOXCENTER, USPATFULL

(\*File contains numerically searchable property data)

# Absolute stereochemistry.

PAGE 1-B

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

123 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

123 REFERENCES IN FILE CAPLUS (1967 TO DATE)

18 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 137:16915

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REFERENCE
            2:
                136:319650
REFERENCE
                136:257619
            3:
REFERENCE
                136:151393
            4:
REFERENCE
            5:
                136:144657
REFERENCE
                136:113039
            6:
REFERENCE
            7:
                136:50169
REFERENCE
                135:253549
            8:
REFERENCE
            9:
                135:192374
REFERENCE
          10:
                135:44718
L141 ANSWER 75 OF 81 REGISTRY COPYRIGHT 2002 ACS
     491-97-4 REGISTRY
     Thymidine 5'-(trihydrogen diphosphate) (9CI)
CN
                                                     (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN
     Thymidine 5'-(trihydrogen pyrophosphate) (8CI)
CN
     Thymidine 5'-pyrophosphate (7CI)
CN
     Thymidine pyrophosphate (6CI)
OTHER NAMES:
CN
     5'-TDP
CN
     dTDP
CN
     TDP
CN
     TDP (nucleotide)
CN
     Thymidine 5'-diphosphate
CN
     Thymidine diphosphate
CN
     Thymidine, mono(trihydrogen diphosphate) (ester)
FS
     STEREOSEARCH
MF
     C10 H16 N2 O11 P2
CI
     COM
LC
     STN Files:
                  AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
       BIOTECHNO, CA, CAOLD, CAPLUS, CASREACT, CIN, CSCHEM, DDFU, DRUGU,
       EMBASE, IFICDB, IFIPAT, IFIUDB, MEDLINE, TOXCENTER, USPATFULL
         (*File contains numerically searchable property data)
```

Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

- 281 REFERENCES IN FILE CA (1967 TO DATE)
- 12 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
- 282 REFERENCES IN FILE CAPLUS (1967 TO DATE)

# 35 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:382185 REFERENCE 136:259178 2: REFERENCE 136:228721 3: REFERENCE 136:114711 4: 136:84678 REFERENCE 5: REFERENCE 6: 136:50169 136:36563 REFERENCE 7: 135:328754 REFERENCE 8: REFERENCE 135:314428 9: REFERENCE 10: 135:303873 L141 ANSWER 76 OF 81 REGISTRY COPYRIGHT 2002 ACS **65-47-4** REGISTRY Cytidine 5'-(tetrahydrogen triphosphate) (8CI, 9CI) (CA INDEX NAME) OTHER NAMES: CN 5'-CTP CN CTP CN Cytidine 5'-triphosphate CN Cytidine triphosphate Cytidine, mono(tetrahydrogen triphosphate) (ester) CN FS STEREOSEARCH MF C9 H16 N3 O14 P3 CI COM AGRICOLA, ANABSTR, BEILSTEIN\*, BIOBUSINESS, BIOSIS, LC

C STN Files: AGRICOLA, ANABSTR, BEILSTEIN\*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CSCHEM, DDFU, DRUGU, EMBASE, IPA, MEDLINE, RTECS\*, TOXCENTER, USPATFULL

(\*File contains numerically searchable property data)
her Sources: EINECS\*\*, NDSL\*\*, TSCA\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

# Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

2537 REFERENCES IN FILE CA (1967 TO DATE)
56 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
2542 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967).

REFERENCE 1: 137:16961

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REFERENCE
                137:2403
            2:
REFERENCE
            3:
                136:397775
REFERENCE
                136:382049
            4:
REFERENCE
                136:365598
            5:
REFERENCE
            6:
                136:352311
REFERENCE
            7:
                136:351991
                136:350567
REFERENCE
            8:
REFERENCE
            9:
                136:336939
REFERENCE
          10:
                136:290862
L141 ANSWER 77 OF 81 REGISTRY COPYRIGHT 2002 ACS
     63-39-8 REGISTRY
     Uridine 5'-(tetrahydrogen triphosphate) (8CI, 9CI)
                                                         (CA INDEX NAME)
CN
OTHER NAMES:
     5'-UTP
CN
     Uridine 5'-triphosphate
CN
CN
     Uridine triphosphate
CN
     Uridine, mono(tetrahydrogen triphosphate) (ester)
CN
     Uteplex
CN
     UTP
FS
     STEREOSEARCH
     C9 H15 N2 O15 P3
MF
CI
     COM
LC
                 ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT,
       CEN, CHEMLIST, CIN, CSCHEM, DDFU, DRUGNL, DRUGU, DRUGUPDATES, EMBASE,
       GMELIN*, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, NIOSHTIC, PROMT,
       RTECS*, TOXCENTER, USPATFULL
         (*File contains numerically searchable property data)
     Other Sources:
                      EINECS**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3572 REFERENCES IN FILE CA (1967 TO DATE)

75 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

3576 REFERENCES IN FILE CAPLUS (1967 TO DATE) 9 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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REFERENCE
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                137:29993
REFERENCE
            2:
                137:28493
REFERENCE
            3:
                137:17751
REFERENCE
            4:
                137:17586
REFERENCE
            5:
                137:17316
                137:16961
REFERENCE
            6:
REFERENCE
            7:
                137:16011
REFERENCE
            8:
                137:5048
REFERENCE
            9:
                137:4461
REFERENCE
          10:
                137:880
L141 ANSWER 78 OF 81 REGISTRY COPYRIGHT 2002 ACS
     63-38-7 REGISTRY
     Cytidine 5'-(trihydrogen diphosphate) (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Cytidine 5'-(trihydrogen pyrophosphate) (8CI)
CN
CN
     Cytidine pyrophosphate (6CI)
OTHER NAMES:
     5'-CDP
CN
CN
     CDP
CN
     Cytidine 5'-diphosphate
CN
     Cytidine 5'-pyrophosphate
     Cytidine coenzyme
CN
CN
     Cytidine diphosphate
CN
     Cytidine, mono(trihydrogen diphosphate) (ester)
FS
     STEREOSEARCH
DR
     87691-21-2
     C9 H15 N3 O11 P2
MF
CI
     COM
LC
                AGRICOLA, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA,
       CANCERLIT, CAOLD, CAPLUS, CASREACT, CEN, CHEMCATS, CHEMLIST, CSCHEM,
       DDFU, DRUGU, EMBASE, GMELIN*, IFICDB, IFIPAT, IFIUDB, MEDLINE, MSDS-OHS,
       RTECS*, TOXCENTER, USPATFULL
         (*File contains numerically searchable property data)
     Other Sources:
                      EINECS**
         (**Enter CHEMLIST File for up-to-date regulatory information)
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<sup>\*\*</sup>PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

964 REFERENCES IN FILE CA (1967 TO DATE)

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145 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
             964 REFERENCES IN FILE CAPLUS (1967 TO DATE)
              67 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
            1: 136:382185
REFERENCE
                136:365598
REFERENCE
            2:
REFERENCE
            3:
                136:350567
REFERENCE
            4:
                136:213828
REFERENCE
            5:
                136:212678
REFERENCE
            6:
                136:84678
REFERENCE
            7:
                136:50169
REFERENCE
            8:
                136:36563
REFERENCE
            9:
                136:17167
REFERENCE
          10:
                136:4799
L141 ANSWER 79 OF 81 REGISTRY COPYRIGHT 2002 ACS
     58-98-0 REGISTRY
     Uridine 5'-(trihydrogen diphosphate) (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Uridine 5'-(trihydrogen pyrophosphate) (8CI)
     Uridine diphosphate (6CI)
     Uridine pyrophosphate (7CI)
CN
OTHER NAMES:
CN
     5'-UDP
CN
CN
     Uridine 5'-diphosphate
     Uridine 5'-pyrophosphate
CN
     Uridine 5'-pyrophosphoric acid
CN
CN
     Uridine, 5'-(trihydrogen diphosphate)
FS
     STEREOSEARCH
     489-66-7, 141342-82-7, 88756-92-7
DR
     C9 H14 N2 O12 P2
MF
CI
     COM
                  ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
LC
     STN Files:
       BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CEN, CHEMLIST,
       CSCHEM, DDFU, DRUGU, EMBASE, GMELIN*, IFICDB, IFIPAT, IFIUDB, MEDLINE,
       MRCK*, NAPRALERT, TOXCENTER, USPATFULL
         (*File contains numerically searchable property data)
                      EINECS**
     Other Sources:
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1483 REFERENCES IN FILE CA (1967 TO DATE)

98 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

1485 REFERENCES IN FILE CAPLUS (1967 TO DATE)

4 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 137:29993

REFERENCE 2: 137:17586

REFERENCE 3: 137:17316

REFERENCE 4: 137:5048

REFERENCE 5: 137:880

REFERENCE 6: 137:400

REFERENCE 7: 136:382185

REFERENCE 8: 136:380468

REFERENCE 9: 136:364761

REFERENCE 10: 136:350567

L141 ANSWER 80 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN **58-64-0** REGISTRY

CN Adenosine 5'-(trihydrogen diphosphate) (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Adenosine 5'-(trihydrogen pyrophosphate) (8CI)

CN Adenosine diphosphate (6CI)

OTHER NAMES:

CN .alpha.-ADP

CN 5'-ADP

CN Adenosine 5'-diphosphate

CN Adenosine 5'-diphosphoric acid

CN Adenosine 5'-pyrophosphate

CN Adenosine 5'-pyrophosphoric acid

CN Adenosine pyrophosphate

CN Adenosine, 5'-(trihydrogen diphosphate)

CN ADP

CN ADP (nucleotide)

FS STEREOSEARCH

DR 84412-16-8

MF C10 H15 N5 O10 P2

CI COM

LC STN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN\*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, DDFU, DETHERM\*, DRUGU, EMBASE, GMELIN\*, IFICDB,

IFIPAT, IFIUDB, IPA, MEDLINE, MRCK\*, NIOSHTIC, PIRA, PROMT, RTECS\*, TOXCENTER, USPATFULL, VETU

(\*File contains numerically searchable property data)

Other Sources: DSL\*\*, EINECS\*\*, TSCA\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

# Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

20334 REFERENCES IN FILE CA (1967 TO DATE)

465 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

20356 REFERENCES IN FILE CAPLUS (1967 TO DATE)

22 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 137:37738

REFERENCE 2: 137:32213

REFERENCE 3: 137:32110

REFERENCE 4: 137:31996

REFERENCE 5: 137:29818

REFERENCE 6: 137:29801

REFERENCE 7: 137:29764

REFERENCE 8: 137:29618

REFERENCE 9: 137:28528

REFERENCE 10: 137:27969

# L141 ANSWER 81 OF 81 REGISTRY COPYRIGHT 2002 ACS

RN 56-65-5 REGISTRY

CN Adenosine 5'-(tetrahydrogen triphosphate) (8CI, 9CI) (CA INDEX NAME) OTHER NAMES:

CN 5'-ATP

CN Adenosine 5'-triphosphate

CN Adenosine 5'-triphosphoric acid

CN Adenosine triphosphate

CN Adenosine, 5'-(tetrahydrogen triphosphate)

CN Adenylpyrophosphoric acid

CN Adephos

CN Adetol

CN Adynol

CN Atipi

CN ATP

```
ATP (nucleotide)
CN
CN
     Atriphos
CN
     Cardenosine
CN
     Fosfobion
CN
     Glucobasin
     Myotriphos
CN
     Phosphobion
CN
CN
     Striadyne
CN
     Triadenyl
CN
     Triphosphaden
CN
     Triphosphoric acid adenosine ester
FS
     STEREOSEARCH
     10168-83-9, 16488-07-6, 51569-41-6, 71800-44-7, 84412-18-0
DR
MF
     C10 H16 N5 O13 P3
CI
     COM
                  ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
LC
     STN Files:
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS,
       CASREACT, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, DDFU, DETHERM*, DRUGNL,
       DRUGU, DRUGUPDATES, EMBASE, GMELIN*, IFICDB, IFIPAT, IFIUDB, IPA,
       MEDLINE, MRCK*, NAPRALERT, NIOSHTIC, PHAR, PIRA, PROMT, RTECS*,
       SPECINFO, TOXCENTER, TULSA, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
                      DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

## Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

60191 REFERENCES IN FILE CA (1967 TO DATE)
1109 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
60244 REFERENCES IN FILE CAPLUS (1967 TO DATE)
19 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 137:37738 REFERENCE 137:32580 REFERENCE 137:32490 REFERENCE 4: 137:32213 REFERENCE 5: 137:32110 137:32036 REFERENCE 6: 137:31996 REFERENCE 7:

8:

137:31532

REFERENCE

REFERENCE 9: 137:31297

REFERENCE 10: 137:31026

=> fil hcaplus

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FILE COVERS 1907 - 16 Jul 2002 VOL 137 ISS 3 FILE LAST UPDATED: 15 Jul 2002 (20020715/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

=> d bib abs hitrn retable tot 1130

L130 ANSWER 1 OF 31 HCAPLUS COPYRIGHT 2002 ACS

AN 2002:332677 HCAPLUS

DN 136:350567

- TI Method of treating gastrointestinal tract disease with purinergic P2Y receptor agonists
- IN Yerxa, Benjamin R.; Rideout, Janet L.; Pendergast, William; Shaver, Sammy R.; Zhang, Zhen; Peterson, Ward M.; Cowlen, Matthew

PA USA

SO U.S. Pat. Appl. Publ., 29 pp., Cont.-in-part of U.S. Ser. No. 512,867. CODEN: USXXCO

DT Patent

LA English

FAN.CNT 3

		TENT NO.	KIND	DATE	APPLICATION NO. DA	DATE		
ΡI		2002052336	A1	20020502		0001222		
	US	6331529	В1	20011218	US 2000-512867 20	0000225		
	US	2002052338	A1	20020502	US 2001-5267 20	0011203		
PRAI	US	1999-121754P	P	19990226				
	US	1999-171710P	P	19991222				
	US	2000-512867	A2	20000225				
			_					

OS MARPAT 136:350567

AB The invention provides a method of regulating water and mucin secretions and fluid transport in the gastrointestinal tract. The invention also provides a method for treating a gastrointestinal disease in which the mucosal barrier of the gastrointestinal system is impaired. The invention addnl. provides a method for correcting disorders of fluid secretion or absorption in the gastrointestinal system. The method comprises

administering to a patient a pharmaceutical compn. comprising a purinergic P2Y receptor agonist, in an amt. effective to regulate water and mucin secretions or to correct abnormal fluid transport in the gastrointestinal tract. The pharmaceutical compn. used in this invention comprises a P2Y purinergic receptor agonist such as UDP, UTP, CDP, CTP, ADP, ATP, and their analogs, as well as dinucleotide polyphosphate compds. The compd. is prepd. in an oral form, an injectable form, or a suppository form, and administered to a patient. IT **34983-48-7**, 2-MethylthioADP RL: PAC (Pharmacological activity); BIOL (Biological study) (purinergic P2Y agonist for treatment of gastrointestinal disease) 56-65-5, Adenosine triphosphate, biological studies ΙT 58-64-0, ADP, biological studies 58-98-0, UDP, biological studies 63-38-7, CDP 63-39-8, UTP 65-47-4, CTP 5542-28-9 5959-90-0 30632-06-5 59985-21-6 63785-59-1 81534-69-2 111648-11-4 211448-70-3 211448-78-1 211448-85-0 345950-34-7 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (purinergic P2Y agonist for treatment of gastrointestinal disease) L130 ANSWER 2 OF 31 HCAPLUS COPYRIGHT 2002 ACS ΑN 2002:256037 HCAPLUS DN 136:273216 Catecholamine adrenergic pharmaceutical compositions IN Root-Bernstein, Robert S.; Dillon, Patrick F. Board of Trustees Operating Michigan State University, USA PA PCT Int. Appl., 40 pp. CODEN: PIXXD2 Patent English LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE \_\_\_\_\_ WO 2002026223 A2 20020404 WO 2001-US30272 20010927 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRAI US 2000-236751P Ρ 20000929 Pharmaceutical compns. comprise (a) a safe and effective amt. of an adrenergic compd.; and (b) a complement to said adrenergic compd., selected from the group consisting of a hyperpreserving amt. of an ascorbate, a safe and effective amt. of an opioid, a hyperpreserving amt. of a polycarboxylic acid chelator, a safe and effective amt. of D-ribose and adenosine derivs., and mixt. thereof. Methods are also provided for regulating an adrenergic receptor in a human or other animal, comprising the administration of: (c) a low dose of an adrenergic compd.; and (d) a safe and effective amt. of a complement to said adrenergic compd. Preferably, the adrenergic compd. is a catecholamine. Preferred complements include ascorbates, particularly ascorbic acid. Methods include the treatment of neurol. disorders, hypotension, forward failure,

backward failure, congestive heart failure, shock, hypertension, hemorrhage, disorders assocd. with anesthesia, chronic obstructive pulmonary disease, asthma, colic, Crohn's disease, anaphylaxis, interstitial cystitis, overactive bladder syndrome, premature labor,

myasthenia gravis, and glaucoma. Asthmatic patients were

ΤI

SO

DT

PΙ

administered an aerosol comprising 0.075% isoproterenol and 1.0% ascorbic acid. Systemic uptake of isoproterenol was decreased, thereby eliminating the adverse side effects previously experienced.

IT 56-65-5, Atp, biological studies

RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(catecholamine adrenergic pharmaceutical compns.)

```
L130 ANSWER 3 OF 31 HCAPLUS COPYRIGHT 2002 ACS
```

AN 2002:136070 HCAPLUS

DN 136:151393

TI Preparation of dinucleotides and their use as modulators of mucociliary clearance and ciliary beat frequency

IN Pendergast, William; Yerxa, Benjamin R.; Rideout, Janet L.; Siddiqi, Suhaib M.

PA Inspire Pharmaceuticals, Inc., USA

SO U.S., 15 pp., Cont.-in-part of U.S. 5,900,407. CODEN: USXXAM

DT Patent

LA English

FAN.CNT 4

	PATENT NO.		KI	ND	DATE			A	PPLI	CATI	N NC	0.	DATE					
ΡΙ			A A	2	20020219 19990504 19981117 19980813 20000106			US 1997-798508			2 8	19980710 19970206 19970210 19980206						
	W:	DK, KP, NO, UA, GH, FR,	EE, KR, NZ, UG, GM, GB,	ES, KZ, PL, US, KE, GR,	FI, LC, PT, US, LS, IE,	AZ, GB, LK, RO, UZ, MW, IT, NE,	GE, LR, RU, VN, SD, LU,	GH, LS, SD, YU, SZ, MC,	GM, LT, SE, ZW, UG, NL,	GW, LU, SG, AM, ZW,	HU, LV, SI, AZ, AT,	ID, MD, SK, BY, BE,	IL, MG, SL, KG, CH,	IS, MK, TJ, KZ, DE,	JP, MN, TM, MD, DK,	KE, MW, TR, RU, ES,	KG, MX, TT, TJ, FI,	TM
PRAI OS GI	US 2002 US 1997 US 1998 WO 1998 US 1998 MARPAT	7-797 7-798 3-US2 3-101	472 508 702 395	A A W A	2 2 ~	2002 1997 1997 1998 1998	0206 <b>0210</b> 0206		U:	S 20	01-7	451		2001	1106			

AB The present invention relates to certain novel dinucleotides I (X = O, CH2, imido, CF2; B, B1 = independently nucleobase; Z, Z1 = independently OH, N3; Y, Y1 = independently H, OH; Q = (HPO3)m; n = 0-2; m = 0-2; n + m = 0-4) and formulations thereof which are highly selective agonists of the P2Y2 and/or P2Y4 purinergic receptor. They are useful in the treatment of chronic obstructive pulmonary diseases such as chronic bronchitis, PCD, cystic fibrosis, as well as prevention of pneumonia due to immobility. Furthermore, because of their general ability to clear retained mucus

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secretions and stimulate ciliary beat frequency, the compds. of the
    present invention are also useful in the treatment of sinusitis, otitis
    media and nasolacrimal duct obstruction. They are also useful for
    treatment of dry eye disease and retinal detachment.
    Thus, P1, P2-di(uridine-5'-)-P2, P3-methylenetetraphosphate was prepd. as
    P2Y2 and/or P2Y4 purinergic receptor (EC50 = 11.1 .mu.mol).
IT
    2596-55-6P 4130-19-2P 5542-28-9P
    5959-90-0P 6674-45-9P 10527-46-5P
    10527-48-7P 13457-68-6P 26184-65-6P
    30632-08-7P 34692-44-9P 41708-91-2P
    56983-23-4P 59985-20-5P 59985-21-6P
    61340-12-3P 63785-59-1P 79695-24-2P
    79695-25-3P 83008-69-9P 88109-92-6P
    96920-51-3P 97776-54-0P 97776-55-1P
    103137-88-8P 103137-89-9P 111035-55-3P
    111648-11-4P 134311-47-0P 135780-83-5P
    135780-85-7P 135780-92-6P 135802-64-1P
    154960-70-0P 170638-56-9P 170638-57-0P
    170638-58-1P 170638-59-2P 170638-60-5P
    170638-61-6P 170638-62-7P 211427-06-4P
    211427-09-7P 211427-11-1P 211448-67-8P
    211448-72-5P 211448-73-6P 211448-74-7P
    211448-76-9P 211448-77-0P 211448-78-1P
    211448-79-2P 211448-80-5P 211448-81-6P
    211448-88-3P
    RL: BSU (Biological study, unclassified); SPN (Synthetic preparation);
    THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (prepn. of dinucleotides and their use as modulators of mucociliary
       clearance and ciliary beat frequency)
IT
    63-39-8, Uridine 5'-triphosphate 19817-92-6
    211448-71-4
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (prepn. of dinucleotides and their use as modulators of mucociliary
       clearance and ciliary beat frequency)
    211427-07-5P 211448-70-3P 211448-75-8P
TΤ
    211448-82-7P 211448-83-8P 211448-84-9P
    211448-85-0P
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (prepn. of dinucleotides and their use as modulators of mucociliary
       clearance and ciliary beat frequency)
RETABLE
  Referenced Author | Year | VOL | PG | Referenced Work
                                                            | Referenced
                     |(RPY)|(RVL)|(RPG)| (RWK)
       (RAU)
                                                            | File
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                      |1998 |
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|1986 |261 |16410 |J Biol Chem

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|Br J Pharmacol

|Br J Pharmacol

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·				Am J Respir Cell Mol	•
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		205		Eur J Biochem	HCAPLUS
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                                 |14009 |Biochemistry | | HCAPLUS
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L130 ANSWER 4 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    2002:107834 HCAPLUS
ΑN
DN
    136:146442
TI
    Method and composition for the accelerated in vivo removal of ethanol
    Bowen; Ward Beryl; Daniel, Daniel Salman
IN
PA
SO
    U.S. Pat. Appl. Publ., 10 pp.
    CODEN: USXXCO
DT
    Patent
    English
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
                                         _____
    ______
                    ____
    US 2002015741 A1
                          20020207
                                         US 2001-876322
                                                          20010607
PI
PRAI US 2000-210950P P 20000612
    A compn. for accelerating the disposal of ethanol from bodily fluid.
    Certain additives can accelerate the metabolic oxidn. of ethanol, and
    others in addn. act as catalysts or "pseudo" enzymes for the oxidn.
    Additives include the oxidant NAD and a variety of other additives such as
    transition metal ions and complexes thereof which favor the oxidn.
    reaction. The compns. described can act as a sobriety inducer and/or as
    an effective palliative for the unpleasant effects of overuse of ethanol.
    56-65-5, 5'-ATP, biological studies
ΙT
    RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (accelerant; method and compn. for accelerated in vivo removal of
       ethanol)
L130 ANSWER 5 OF 31 HCAPLUS COPYRIGHT 2002 ACS
ΑN
    2002:10323 HCAPLUS
DN
    136:74708
    Composition and method for the repair and regeneration of cartilage and
ΤI
    other tissues based on a polymer gel
    Hoemann, Caroline D.; Buschmann, Michael D.; Mckee, Marc D.
ΙN
    Biosyntech Canada Inc., Can.
PΑ
SO
    PCT Int. Appl., 106 pp.
    CODEN: PIXXD2
DT
    Patent
    English
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                        APPLICATION NO. DATE
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                                         _____
                                    WO 2001-CA959 20010629
                          20020103
    WO 2002000272
                    A2
PΙ
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT,
            RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US,
            UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,

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DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     AU 2001068882
                            20020108
                                           AU 2001-68882
                                                            20010629
                       Α5
     US 2002082220
                                           US 2001-896912
                                                            20010629
                       Α1
                            20020627
PRAI US 2000-214717P
                       Ρ
                            20000629
     WO 2001-CA959
                       W
                            20010629
     The present invention relates to a new method for repairing human or
AB
     animal tissues such as cartilage, meniscus, ligament, tendon, bone, skin,
     cornea, periodontal tissues, abscesses, resected tumors, and ulcers. The
     method comprises the step of introducing into the tissue a temp.-dependent
     polymer gel compn. such that the compn. adhere to the tissue and promote
     support for cell proliferation for repairing the tissue. Other than a
     polymer, the compn. preferably comprises a blood component such as whole
     blood, processed blood, venous blood, arterial blood, blood from bone,
     blood from bone-marrow, bone marrow, umbilical cord blood, placenta blood,
     erythrocytes, leukocytes, monocytes, platelets, fibrinogen, thrombin and
     platelet rich plasma. The present invention also relates to a new compn.
     to be used with the method of the present invention. For example,
     chondral defects with perforations to the subchrondal bone of rabbits were
     treated with a peripheral blood/chitosan-glyceryl phosphate mixt. that was
     delivered as a liq., and allowed to solidify in situ. After 5-8 wk
     healing, the blood/chitosan-treated defects were filled with repair tissue
     having the appearance of hyaline, a glycosaminoglycan (GAG)-rich cartilage
     repair tissue, which adhered to the defect surfaces, and filled the
                Repair tissue from the untreated defects (control) had the
     appearance of fibro-cartilage, with particularly no metachromatic staining
     for GAG, and only partial defect filling.
     58-64-0, Adenosine diphosphate, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (temp.-dependent polymer gel compns. contg. blood components for repair
        and regeneration of human or animal tissues)
L130 ANSWER 6 OF 31 HCAPLUS COPYRIGHT 2002 ACS
AN
     2001:885819 HCAPLUS
DN
     136:705
     Ectocornea extension promoters containing P2Y receptor agonists
TI
     Nakata, Katsuhiko; Nakamura, Masatsugu; Fujihara, Tsutomu; Fujita, Hiromi
ΙN
     Santen Pharmaceutical Co., Ltd., Japan; Inspire Pharmaceuticals, Inc.
PA
SO
     PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
DT
     Patent
T.A
     Japanese
FAN.CNT 1
                                           APPLICATION NO. DATE
     PATENT NO.
                      KIND
                            DATE
     WO 2001091795
                      A1
                            20011206
                                           WO 2001-JP4520
                                                            20010530
ΡI
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
             HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
             LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD,
             SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU,
             ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                            20020219
     JP 2002053492
                      A2
                                           JP 2001-161664
                            20000530
PRAI JP 2000-159889
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AB As the results of searching for compds. having an effect of promoting ectocornea extension in the ophthalmol. field, it is found out that P2Y receptor agonists typified by phosphoric acid compds. having adenosyl, uridyl, xanthosyl, guanosyl or thymidyl and salts thereof exhibit an excellent effect of promoting ectocornea extension. The effect of

MARPAT 136:705

OS

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P1,P4-di(uridine-5')-tetraphosphate tetrasodium salt (DUTP-Na) on rabbit
    corneal epithelium elongation was examd. Also, an eye drop contg. DUTP-Na
    10, NaCl 90 mg, and water q.s. to 100 mL was formulated.
    16178-48-6, Adenosine 5'-diphosphate disodium salt
ΙT
    19817-92-6, Uridine 5'-triphosphate trisodium salt
    27821-45-0, Uridine 5'-diphosphate disodium salt
    211427-08-6
    RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (P2Y receptor agonists as corneal epithelium elongation promoters)
RETABLE
  Referenced Author | Year | VOL | PG | Referenced Work
                                                            | Referenced
                    | (RPY) | (RVL) | (RPG) | (RWK)
                                                           | File
        (RAU)
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                                      |EP 1087777 A2
Inspire Pharmaceuticals
                         1
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Inspire Pharmaceuticals
                                       IUS 5900407 A
                                                          | HCAPLUS
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Inspire Pharmaceuticals
                                                          HCAPLUS
Inspire Pharmaceuticals
                                      IAU 9863242 A1
                                                           HCAPLUS
Inspire Pharmaceuticals
                                       |AU 9940839 A
                                                           | HCAPLUS
                                       IWO 9834942 A2
Inspire Pharmaceuticals | 1998 |
                                                           | HCAPLUS
Inspire Pharmaceuticals | 1999 |
                                        WO 9961012 A2
                                                           HCAPLUS
                                        IWO 0050024 A2
Inspire Pharmaceuticals|2000 |
                                                            | HCAPLUS
                                 |129
                                        |ARCHIVES OF HISTOLOG|HCAPLUS
                      |1999 |62
Kimura, K
L130 ANSWER 7 OF 31 HCAPLUS COPYRIGHT 2002 ACS
ΑN
    2001:851183 HCAPLUS
DN
    136:690
ΤI
    Method for retinal degeneration treatment with purinergic
    receptor agonists
IN
    Peterson, Ward M.
    Inspire Pharmaceuticals, Inc., USA
PA
SO
    PCT Int. Appl., 46 pp.
    CODEN: PIXXD2
DT
    Patent
    English
LA
FAN.CNT 1
                   KIND DATE
                                       APPLICATION NO. DATE
    PATENT NO.
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                                         _____
                                                         _____
    WO 2001087913
                  A2
                          20011122
                                         WO 2001-US15606 20010510
PΙ
    WO 2001087913
                    A3
                          20020530
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
        W:
            CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
            HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
            LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO,
            RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ,
            VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRAI US 2000-570231
                          20000512
                     Α
OS
    MARPAT 136:690
AΒ
    Methods are disclosed for prevention or treatment of retinal
    degeneration arising from pathophysiol. or phys. conditions. The method
    involves administration of a pharmaceutical compn. comprising a purinergic
    P2Y receptor ligand, in an amt. effective to elevate its extracellular
    concn. to activate retinal glial and neuronal cell surface P2Y
    receptors and mount a neuroprotective response. Also disclosed are
    methods of administration including intravitreal bolus and
    sustained-release administration, transscleral delivery, topical, and
    systemic administration. The pharmaceutical compn. useful in the
    invention comprises a P2Y purinergic receptor agonist, which include
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uridine 5'-di -and triphosphate (UDP, UTP) and their analogs, ADP (ADP)

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and its analogs, cytidine 5'-di- and triphosphate (CDP, CTP) and their
    analogs, and dinucleoside polyphosphate compds.
ΙT
     63-39-8, Uridine 5'-triphosphate 65-47-4, Cytidine
     5'-triphosphate 5542-28-9, AP4A 5959-90-0
    30632-06-5 59985-21-6 63785-59-1
    81534-69-2 111648-11-4 188560-02-3
    211448-70-3 211448-78-1 211448-85-0
    345950-34-7
    RL: BSU (Biological study, unclassified); THU (Therapeutic use);
    BIOL (Biological study); USES (Uses)
        (method for retinal degeneration treatment with purinergic
        receptor agonists)
L130 ANSWER 8 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    2001:763523 HCAPLUS
AN
DN
    135:298823
TΙ
    Use of P2Y receptor agonist dinucleotide compounds to stimulate removal of
    fluid in retinal detachment and retinal
    Peterson, Ward M.; Yerxa, Benjamin R.
IN
    Peterson, Ward M., USA
PΑ
    U.S. Pat. Appl. Publ., 20 pp., Cont.-in-part of U.S. 5,837,861.
SO
    CODEN: USXXCO
DΨ
    Patent
LA
    English
FAN.CNT 4
    PATENT NO.
                     KIND DATE
                                          APPLICATION NO.
                                                           DATE
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                                          US 2001-774752
                     A1
                           20011018
                                                           20010130
    US 2001031743
PΙ
                      Α
                                          US 1997-798508
                                                           19970210
    US 5837861
                           19981117
                     Α
                           19990219
                                          ZA 1998-1073
                                                           19980210
    ZA 9801073
                    A2
PRAI US 1997-798508
                           19970210
OS
    MARPAT 135:298823
    The invention provides a method of treating edematous retinal
AΒ
    disorders. The method comprises administration of a P2Y receptor agonist
    to stimulate the removal of pathol. extraneous fluid from the subretinal
    and retinal spaces and thereby reduce the accumulation of said
    fluid assocd. with retinal detachment and
    retinal edema. The P2Y receptor agonist may be administered with
    therapeutic and adjuvant agents commonly used to treat edematous
    retinal disorders. The pharmaceutical compn. useful in this
    invention comprises a P2Y receptor agonist with enhanced resistance to
    extracellular hydrolysis, such as dinucleoside polyphosphate compds.
    5542-28-9 10527-48-7 59985-21-6
IT
    83008-69-9 96920-51-3 111648-11-4
    148503-84-8 211427-06-4 211427-07-5
    211427-09-7 211427-10-0 211427-11-1
    211448-70-3 211448-72-5 211448-73-6
    211448-74-7 211448-75-8 211448-76-9
    211448-77-0-211448-78-1-211448-79-2-
    211448-80-5 211448-81-6 211448-82-7
    211448-83-8 211448-84-9 211448-85-0
    318250-11-2, INS 37217 366004-15-1 366004-16-2
    366004-17-3 366004-18-4 366004-19-5
    366004-20-8 366004-21-9 366004-22-0
    366004-23-1
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
    study, unclassified); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (P2Y receptor agonist dinucleotide compds. to stimulate removal of
        fluid in retinal detachment and retinal
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edema)

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L130 ANSWER 9 OF 31 HCAPLUS COPYRIGHT 2002 ACS
AN
     2001:472479 HCAPLUS
     135:71289
DN
TI
    Method of treating gastrointestinal tract disease with purinergic receptor
     agonists
IN
     Yerxa, Benjamin R.; Rideout, Janet L.; Pendergast, William; Shaver, Sammy
     R.; Zhang, Zhen; Peterson, Ward M.; Cowlen, Mathew
PA
     Inspire Pharmaceuticals, Inc., USA
     PCT Int. Appl., 58 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
    English
FAN.CNT 3
                     KIND DATE
                                                            DATE
     PATENT NO.
                                          APPLICATION NO.
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                                          -----
    WO 2001045691
                      A2
                            20010628
                                          WO 2000-US35439 20001222
ΡI
    WO 2001045691
                      A3
                            20020418
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
            HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
            LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
             YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRAI US 1999-171710P
                            19991222
OS
    MARPAT 135:71289
    The invention provides a method of regulating water and mucin secretions
AΒ
    and fluid transport in the gastrointestinal tract. The invention also
    provides a method for treating a gastrointestinal disease in which the
    mucosal barrier of the gastrointestinal system is impaired. The invention
    addnl. provides a method for correcting disorders of fluid secretion or
    absorption in the gastrointestinal system. The method comprises
    administering to a patient a pharmaceutical compn. comprising a purinergic
    P2Y receptor agonist, in an amt. effective to regulate water and mucin
    secretions or to correct abnormal fluid transport in the gastrointestinal
            The pharmaceutical compn. used in this invention comprises a P2Y
    purinergic receptor agonist such as UDP, UTP, CDP, CTP, ADP, ATP, and
    their analogs, as well as dinucleotide polyphosphate compds. The compd.
    is prepd. in an oral, injectable, or suppository form, and administered to
    a patient.
    56-65-5, Adenosine 5'-triphosphate, biological studies
    58-64-0, Adenosine 5'-diphosphate, biological studies
    58-98-0, Uridine 5'-diphosphate, biological studies
    63-38-7, Cytidine 5'-diphosphate 63-39-8, Uridine
    5'-triphosphate 65-47-4, Cytidine 5'-triphosphate
    5542-28-9 5959-90-0 30632-06-5
    34983-48-7, 2-Methylthio-ADP 59985-21-6
    63785-59-1 81534-69-2 111648-11-4
    211448-70-3 211448-78-1 211448-85-0
    345950-34-7
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
    study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (purinergic P2Y receptor agonists for treating gastrointestinal
        disease)
L130 ANSWER 10 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    2001:342036 HCAPLUS
AN
DN
     135:147383
ΤI
    Therapeutic potential of nucleotides in the eye
```

Pintor, Jesus; Peral, Assumpta

- CS Departamento de Bioquimica, E.U. Optica, Universidad Complutense de Madrid, Madrid, 28037, Spain
- SO Drug Development Research (2001), 52(1/2), 190-195 CODEN: DDREDK; ISSN: 0272-4391
- PB Wiley-Liss, Inc.
- DT Journal
- LA English
- Adenine mononucleotides present the ability to produce changes in the intraocular pressure in New Zealand rabbits. Two main groups of compds. were found in terms of their behavior on the one hand, those that increased intraocular pressure, and on the other hand, those that reduced it. The hypotensive compds. showed a concn.-response behavior, and were blocked by the P2 receptor antagonist pyridoxalphosphate-6-azophenyl-2',4'-disulfonic acid. Moreover, the hypotensive compds. were unaffected by the pretreatment with sympathetic antagonists but were completely abolished by the parasympathetic antagonists. This fact suggests the possibility of these P2 receptors to be present at the cholinergic terminal that control intraocular pressure. Adenine mononucleotides also were assayed in their ability to increase tear secretion rate. This study showed that tear secretion is stimulated via P2Y2 receptors.
- IT 56-65-5, Atp, biological studies 58-64-0, Adp,
  biological studies 58-98-0, Udp, biological studies
  63-39-8, Utp
  - RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(therapeutic potential of nucleotides in ophthalmol.)

#### RETABLE

Referenced Author	lYear	I VOL	I PG	Referenced Work	Referenced
(RAU)	(RPY)	•	•	(RWK)	File
				-+=====================================	+========
Abe, Y	11995	64	547	Neuroscience	HCAPLUS
Burnstock, G	11994	31	206	Drug Dev Res	
Burnstock, G	•		67		HCAPLUS
Crosson, C	1996	137	1833	Invest Ophthalmol Vi	MEDLINE
Crosson, C	11994	110	379	J Ocular Pharmacol	HCAPLUS
Crosson, C	1995	1273	320	J Pharmacol Exp Ther	
Davson, H	11993	Ι.	34	Physiology of the ey	
Harden, T	11995	35	541	Ann Rev Pharmacol To	HCAPLUS
Howard, M	11998	139	1942	Invest Ophthalmol Vi	MEDLINE
James, S	11993	160	219		HCAPLUS
Kaufman, P	1984	1	149	Handbook of experime	HCAPLUS
Lambrecht, G	1992	217	217	Eur J Pharmacol	HCAPLUS
Lutjen-Drecoll, J	1989	1	189	The glaucoma	1
Maul, E	1979	18	256	Invest Ophthalmol Vi	HCAPLUS
Nishimura, T	1996	212	215	Neurosci Lett	HCAPLUS
Northway, M	1980	65	11	Eur J Pharmacol	HCAPLUS
Pintor, J	12000	l	171	Nervous control of t	
Ralevic, V	1998	150	413	•	HCAPLUS
Redman, R	1994 -	477	1117-	J Physiol (Lond) -	HCAPLUS-
Richardson, P	1987	48	622	J Neurochem	HCAPLUS
Rohen, J	1982	1	141	Basic aspect of glau	
Rohen, J	1964	1 .	189	Handbuch der mikrosk	i
Shahidullah, M	1997	16	1006	Curr Eye Res	MEDLINE
Sun, X	•	93	1859	Proc Natl Acad Sci U	HCAPLUS
Ueno, S	11992	68	1778		HCAPLUS
Wiederholt, M	1998	1	1163	The eye's aqueous hu	HCAPLUS
Yerxa, B	11999	1	B723	Invest Ophthalmol Vi	1
Zimmemmann, H	11993	1	1	Synaptic transmissio	1
Zimmermann, H	1996	139	1337	Drug Develop Res	HCAPLUS

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DN
    134:141776
    Methods for transdifferentiation of body tissues
ΤI
    Baranowitz, Steven
IN
PA
    USA
    PCT Int. Appl., 51 pp.
SO
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 1
     PATENT NO.
                 KIND DATE
                                 APPLICATION NO. DATE
    WO 2001008691 A1 20010208 WO 2000-US21015 20000731
ΡI
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL,
            IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
            MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG,
            SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW,
            AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
            CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
    EP 1200101
                     A1 20020502 EP 2000-955323 20000731
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL
PRAI US 1999-146272P P 19990729
    US 1999-168555P P
                           19991202
    WO 2000-US21015 W 20000731
AB
    This invention relates to methods for transdifferentiation of body tissues
    which can be used to generate specific cell types needed for regenerating
    organs or body parts, following cellular degeneration, injury or
    amputation. The present invention also describes the use of tissue
    transdifferentiation for treating cancer and autoimmune diseases.
    56-65-5, Adenosine triphosphate, biological studies
IT
    58-64-0, Adenosine diphosphate, biological studies 58-98-0
     , Uridine diphosphate, biological studies 63-39-8, Uridine
    triphosphate 491-97-4, Thymidine diphosphate
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
    study, unclassified); THU (Therapeutic use); BIOL (Biological
    study); USES (Uses)
        (transdifferentiation by; methods for transdifferentiation of body
        tissues for use for regenerating organs or body parts or treating
        cancer and autoimmune disease)
RETABLE
  Referenced Author | Year | VOL | PG | Referenced Work | Referenced
     (RAU) | (RPY) | (RVL) | (RPG) | (RWK)
______________________
Levesque | 2000 | | | US 6087168 A | | HCAPLUS Noji, S | 1991 | 350 | 83 | | Retinoic acid induce | HCAPLUS
                     |1991 |66 |199 |Cell
                                                         | HCAPLUS
Tabin, C
                                      | Cell | HCAPLUS | HCAPLUS | HCAPLUS
Wanek, N | 1991 | 350 | 181
L130 ANSWER 12 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    2000:608596 HCAPLUS
ΑN
DN
    133:187988
    Methods and compositions for altering mucus secretion
ΤI
    Li, Yuehua; Martin, Linda D.; Adler, Kenneth B.
TN
    North Carolina State University, USA
PA
SO
    PCT Int. Appl., 66 pp.
    CODEN: PIXXD2
DT
    Patent
    English
LA
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APPLICATION NO. DATE

FAN.CNT 1

PATENT NO.

KIND DATE

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WO 2000050062
                                                            20000224
PΙ
                       Α2
                            20000831
                                           WO 2000-US5050
     WO 2000050062
                      А3
                            20001221
            AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
             DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
             KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN,
             MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
             TR, TT,
                    UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD,
             RU, TJ,
                    TM
                    KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
         RW: GH, GM,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
                    CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
             CG, CI,
                                                            20000224
                                          EP 2000-912034
     EP 1154786
                      A2
                          20011121
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
PRAI US 1999-256154
                            19990224
                     Α
                            20000224
     WO 2000-US5050
                       W
     Methods and compds. for increasing or decreasing mucus secretion in
AΒ
     subjects, and particularly mucus secretion in the airways, are described.
     The use of compds. that modulate MARCKS protein-related mucus secretion is
     described. Methods of screening compds. for the ability to increase or
     decrease mucus secretion are also described.
     63-39-8, Uridine 5'-triphosphate
ΙT
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (methods and compns. for altering MARCKS protein-related mucus
        secretion)
L130 ANSWER 13 OF 31 HCAPLUS COPYRIGHT 2002 ACS
     2000:608567 HCAPLUS
AN
DN
     133:187963
ΤI
     Method of promoting mucosal hydration with certain uridine, adenine and
     cytidine diphosphates and analogs thereof
     Yerxa, Benjamin R.; Rideout, Janet L.; Jones, Arthur C.
ΙN
     Inspire Pharmaceuticals, Inc., USA
PA
     PCT Int. Appl., 38 pp.
SO
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 3
                      KIND DATE
     PATENT NO.
                                           APPLICATION NO.
                                                            DATE
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                           _____
                                           WO 2000-US5282
                                                             20000225
PΙ
     WO 2000050024
                      A2
                            20000831
     WO 2000050024
                      A3
                            20010705
            AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
             KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN,
             MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
                     UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU,
             TR, TT,
             TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                             20000225
     EP 1161246
                       A2
                           20011212
                                           EP 2000-914781
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
                                           BR 2000-8498
                                                             20000225
     BR 2000008498
                            20020205
                       Α
PRAI US 1999-121754P
                       Ρ
                            19990226
     WO 2000-US5282
                       W
                            20000225
OS
     MARPAT 133:187963
     A method and prepn. for the stimulation of mucosal hydration in a subject
AΒ
     in need of such treatment is disclosed. The method comprises
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administering to the mucosal surfaces of the subject a purinergic receptor agonist such as UDP, dinucleotides, CDP, ADP, or their therapeutically useful analogs and derivs., in an amt. effective to stimulate mucin secretion. Pharmaceutical formulations and methods of making the same are also disclosed. Methods of administering the same would include: topical administration via a liq., gel, cream, or as part of a contact lens or selective release membrane; or systemic administration via nasal drops or spray, inhalation by nebulizer or other device, oral form (liq. or pill), injectable, intra-operative instillation or suppository form. A method for facilitating the expectoration of sputum for the purpose of detecting cellular abnormalities indicative of lung disease is also disclosed. Uridine diphosphate at 104-10-8 M increased mucin release in cultured epithelial and goblets cells.

IT 58-64-0, Adenosine 5'-(trihydrogen diphosphate), biological studies 58-98-0, Uridine diphosphate, biological studies 63-38-7, Cytidine diphosphate

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(method of promoting mucosal hydration with certain uridine, adenine and cytidine diphosphates and analogs thereof)

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L130 ANSWER 14 OF 31 HCAPLUS COPYRIGHT 2002 ACS
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AN 2000:456859 HCAPLUS

DN 133:79356

TI Synthetic and therapeutic methods for the alpha and beta domains of metallothionein

IN Vallee, Bert L.

PA USA

SO PCT Int. Appl., 64 pp. CODEN: PIXXD2

DT Patent

LA English

FAN CNT 1

L WIM .	CIAI	Τ.																
	PAT	ENT	NO.		KIND DATE			APPLICATION NO.				ο.	DATE					
PI	WO 2000038654			54	A1 20000706			WO 1999-US30573 19991221										
		W:	ΑE,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CR,	CU,
			CZ,	DE,	DK,	EE,	ES,	FΙ,	GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,
			IS,	JP,	KE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,
			MG,	MK,	MN,	MW,	MX,	NO,	ΝZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,
			SL,	ТJ,	TM,	TR,	TT,	UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZW,	AM,	ΑZ,	BY,
			KG,	ΚZ,	MD,	RU,	ТJ,	MT										
		RW:	GH,	GM,	ΚE,	LS,	MW,	SD,	SL,	SZ,	TZ,	ÜG,	ZW,	ΑT,	BE,	CH,	CY,	DE,
			DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	ΙΤ,	LU,	MC,	NL,	PΤ,	SE,	BF,	ВJ,	CF,
			CG,	CI,	CM,	GA,	GN,	GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG				

PRAI US 1998-113459P P 19981223

The present invention relates to the alpha and beta domains of metallothionein and analogs thereof, their synthesis, and therapeutic applications of them. Purified metal-free and metal-contg. alpha and beta domains of metallothionein are provided. A high yield method of synthesis and purifn. is also provided for the metal-free and metal-contg. alpha and beta domains of metallothionein. Finally, therapeutic methods are provided that use the alpha and beta domains of metallothionein to transport selected metals to specific tissues or to sequester metals from these tissues in order to treat conditions in those tissues that are ameliorated by the addn. or sequestration of these metals.

IT 56-65-5, 5'-Atp, biological studies

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(synthetic and therapeutic methods for the alpha and beta domains of metallothionein)

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RETABLE
  Referenced Author | Year | VOL | PG | Referenced Work
                                                       | Referenced
       (RAU) | (RPY) | (RVL) | (RPG) | (RWK)
                                                       | File
Bofill Post
Goldenberg
                    |1990 |112 |2291 |Journal of the Ameri|HCAPLUS
Kull
                    |1985 |260 |8698 |The Journal Of Biolo|HCAPLUS
Nielson
                    |1994 |202 |621 |Biochemical And Biop|HCAPLUS
Pan
                    |
|1975 | |
|1988 | |
                                      |US 3910872 A | | HCAPLUS
Riniker
Tolman, G
                                     |US 4732864 A
                                                       HCAPLUS
                    |1988 |
                    |1984 |259 |11419 |The Journal Of Biolo|HCAPLUS
Winge
                    |1979 |76 |486 |Proceedings of the N|HCAPLUS
Yoshida
L130 ANSWER 15 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    2000:393890 HCAPLUS
DN
    133:838
    Method for treating hemorrhage into eye internal media and sclera and
ΤI
ΙN
    Mukha, A. I.
PΑ
    Russia
SO
    Russ.
    From: Izobreteniya 1998, (27), 352.
    CODEN: RUXXE7
DT
    Patent
LA
    Russian
FAN.CNT 1
    PATENT NO.
               KIND DATE
                                     APPLICATION NO. DATE
    _____
                                      _____
                                                     _____
    RU 2119316 C1 19980927
                                     RU 1996-113149 19960624
PΙ
    Title only translated.
AB
    56-65-5, 5'-ATP, biological studies
IT
    RL: ANT (Analyte); BOC (Biological occurrence); BSU (Biological study,
    unclassified); THU (Therapeutic use); ANST (Analytical study);
    BIOL (Biological study); OCCU (Occurrence); USES (Uses)
       (method for treating hemorrhage into eye internal media and sclera and
       retina)
L130 ANSWER 16 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    2000:335023 HCAPLUS
    132:339428
DN
    Defined serum-free medical solution for ophthalmology
TΙ
ΤN
    Skelnik, Debra A.
PA
    Bausch and Lomb Surgical, Inc., USA
SO
    Eur. Pat. Appl., 27 pp.
    CODEN: EPXXDW
DT
    Patent
    English
T.A
FAN.CNT 1
    PATENT NO. KIND DATE APPLICATION NO. DATE
    EP 1000541 A1 20000517 EP 1999-308702 19991102
PΙ
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
           IE, SI, LT, LV, FI, RO
    US 6153582
                                      US 1998-186580
                                                     19981105
                   A
                         20001128
    AU 9957108
                    A1
                         20000511
                                      AU 1999-57108
                                                     19991028
                                      JP 1999-313063 19991102
    JP 2000198701
                   A2
                         20000718
                        19981105
PRAI US 1998-186580
                   Α
    The title soln. contains one or more cell nutrient supplements and a
    growth factor which maintains and enhances the preservation of eye
    tissues, including human corneal, retinal, and corneal
    epithelial tissues at low to physiol. temp. (2-38.degree.). This soln. is
    composed of a defined aq. nutrient and electrolyte soln., supplemented
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with glycosaminoglycans, deturgescent agents, energy sources, buffer systems, antioxidants, membrane stabilizers, antibiotics, antimycotics, ATP or energy precursors, nutrient cell supplements, nonessential amino acids, trace minerals, trace elements, and growth factors. 14264-46-1, Sodium UTP TΤ RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (serum-free medical solns. for ophthalmol.) RETABLE Referenced Author |Year | VOL | PG | Referenced Work Referenced (RAU) | (RPY) | (RVL) | (RPG) | (RWK) | File \_ |EP 0517972 A Lindstrom, R |1992 | |HCAPLUS L130 ANSWER 17 OF 31 HCAPLUS COPYRIGHT 2002 ACS AN 2000:246640 HCAPLUS DN 132:241937 Citicoline compositions for treating optic nerve atrophy ΤI ΙN Wang, Haiyan PA Peop. Rep. China Faming Zhuanli Shenqing Gongkai Shuomingshu, 5 pp. SO CODEN: CNXXEV DT Patent Chinese LA FAN.CNT 1 KIND DATE PATENT NO. APPLICATION NO. DATE CN 1195526 A 19981014 CN 1994-103533 19940421 PΙ The title compns. [injections] comprise citicoline 60, arginine 1, ATP 1, AΒ inosine 30, procaine 0.5, buffer soln. (pH 4-8) 0.5 parts, and injection water 2 mL. The compns. are prepd. by mixing the ingredients, ultrafiltering, drying by freezing, and sterilizing with Co-60. ΙT 56-65-5, ATP, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (citicoline compns. for treating optic nerve atrophy) L130 ANSWER 18 OF 31 HCAPLUS COPYRIGHT 2002 ACS 1999:763861 HCAPLUS ΑN DN Therapeutic dinucleotide and derivatives ΤI IN Yerxa, Benjamin R.; Pendergast, William; Rideout, Janet L.; Picher, Maryse; Boucher, Richard C.; Stutts, Monroe Jackson Inspire Pharmaceuticals, Inc., USA; The University of North Carolina at PA Chapel Hill SO PCT Int. Appl., 23 pp. CODEN: PIXXD2 DT Patent English LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE \_\_\_\_\_ ----WO 9961012 A2 19991202 A3 20000210 WO 1999-US10948 19990519 PΙ WO 9961012 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,

RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,

AU 1999-40839

19990519

CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

19991213

MD, RU, TJ, TM

AU 9940839

A1

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AU 746750
                            20020502
                      В2
    EP 1087777
                      A2
                           20010404
                                          EP 1999-924313
                                                          19990519
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
     JP 2002516273
                      T2
                           20020604
                                          JP 2000-550472
                                                           19990519
    US 6323187
                      В1
                           20011127
                                          US 1999-316571
                                                           19990521
PRAI US 1998-86543P
                      Ρ
                           19980522
    WO 1999-US10948 W
                           19990519
AΒ
    The present invention relates to P1-(cytidine-5'-)-P4-(uridine
     5'-)tetraphosphate and its salts, esters and amides, and formulations
    thereof which are highly stable and selective agonists of the P2Y2 and/or
    P2Y4 purinergic receptor. The compds. of the invention are useful in the
    treatment of chronic obstructive pulmonary diseases such as chronic
    bronchitis, primary ciliary dyskinesia, cystic fibrosis, as well as
    prevention of pneumonia due to immobility, and the induction of sputum and
    its expectoration. Furthermore, because of their general ability to clear
    retained mucus secretions and stimulate ciliary beat frequency, the
    compds. of the present invention are also useful in the treatment of
    sinusitis and otitis media.
TΤ
    251317-44-9P
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
    study, unclassified); SPN (Synthetic preparation); THU (Therapeutic
    use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (dinucleotides as selective purinoceptor agonists for treatment of
        obstructive pulmonary diseases)
    211448-85-0 211448-85-0D, esters
IT
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
    study, unclassified); THU (Therapeutic use); BIOL (Biological
    study); USES (Uses)
        (dinucleotides as selective purinoceptor agonists for treatment of
       obstructive pulmonary diseases)
L130 ANSWER 19 OF 31 HCAPLUS COPYRIGHT 2002 ACS
    1999:388060 HCAPLUS
ΑN
DN
    131:31034
    Purification and therapeutic application of peptide complexes with heat
ΤI
    shock proteins
IN
    Wallen, Erik S.; Moseley, Pope L.
    The University of New Mexico, USA
PA
SO
    PCT Int. Appl., 33 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 3
    PATENT NO.
                     KIND
                           DATE
                                          APPLICATION NO. DATE
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                                         WO 1998-US25734 19981204
    WO 9929182
                           19990617
PΤ
                     A1
        W: BR, CA, JP, MX
        RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT, SE-
    US 5981706
                           19991109
                                          US 1997-986234
                                                           19971205
                      Α
    CA 2312049
                      AΑ
                           19990617
                                          CA 1998-2312049 19981204
                                          EP 1998-961905 19981204
    EP 1035780
                      Α1
                           20000920
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
                      Т2
                                          JP 2000-523867 19981204
    JP 2001525347
                           20011211
PRAI US 1997-985548
                      Α
                           19971205
    US 1997-986234
                      Α
                           19971205
    US 1996-717239
                      Α2
                           19960920
    US 1997-934139
                      Α2
                           19970919
    WO 1998-US25734
                      W
                           19981204
    The authors disclose methods for synthesizing heat shock protein
AΒ
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(hsp)-peptide complexes. The complexes are prepd. by capturing the hsps

on agarose-immobilized gelatin and effecting their elution with the derived peptide(s). Alternatively, the heat shock proteins are captured on an affinity matrix as complexes with ADP prior to their subsequent elution with peptide(s). In addn., the present invention also provides a method for treating an allergic disease in which a heat shock protein-antigen complex is administered to a mammal in an amt. sufficient to reduce the susceptibility of the mammal to a Th2 response for the allergic disease. In an example of desensitization, mice were pretreated with HSP70 complexes contg. peptides derived from the Fel d 1 allergen prior to antigen challenge. ΙT 56-65-5, 5'-ATP, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (for in situ generation of ADP-heat-shock protein complexes) IT 58-64-0DP, 5'-ADP, heat-shock protein-peptide complexes RL: BPN (Biosynthetic preparation); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (prepn., purifn., and antiallergenic application of) RETABLE | Referenced |Year | VOL | PG | Referenced Work Referenced Author |(RPY)|(RVL)|(RPG)| . (RWK) | File (RAU) \_ |16792 |Journal of Biologica| |1996 |278 |1994 |269 |13107 |Journal of Biologica|HCAPLUS Palleros |1997 |204 |13 |Journal of Immunolog|HCAPLUS Peng |1994 |6 1728 |Current Opinions in | HCAPLUS Srivastava |1994 |39 193 |Immunogenetics | HCAPLUS Srivastava Srivastava, P |1993 |62 1153 |Advances in Cancer R|HCAPLUS L130 ANSWER 20 OF 31 HCAPLUS COPYRIGHT 2002 ACS 1999:184143 HCAPLUS AN DN 130:218318 Use of purine nucleosides for modulating the axonal outgrowth of central TΙ nervous system neurons ΤN Benowitz, Larry I. Children's Medical Center Corporation, USA PA PCT Int. Appl., 43 pp. SO CODEN: PIXXD2 DT Patent English LΆ FAN.CNT 1 APPLICATION NO. DATE PATENT NO. KIND DATE ----\_\_\_\_\_ A1 19990311 WO 9911274 WO 1998-US3001 19980220 PΙ W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG CA 2302156 AA19990311 CA 1998-2302156 19980220 AU 9866568 A1 19990322 AU 1998-66568 19980220 20000621 EP 1998-908565 19980220 EP 1009412 A1 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI 20011002 JP 2000-508376 · 19980220 JP 2001516695 T2 20020411 US 2001-997688 20011129 US 2002042390 Α1 US 2001-997687 20011129 US 2002055484 20020509 Α1 PRAI US 1997-921902 Α2 19970902

WO 1998-US3001

W

19980220

owens - 09 / 570231 AB Methods and compns. for modulating the axonal outgrowth of central nervous system neurons are provided. Methods for stimulating the axonal outgrowth of central nervous system neurons following an injury (e.g., stroke, Traumatic Brain Injury, cerebral aneurism, spinal cord injury and the like) and methods for inhibiting the axonal outgrowth of central nervous system neurons in conditions such as epilepsy, e.g., post-traumatic epilepsy, and neuropathic pain syndrome, are also provided. These methods generally involve contacting the central nervous system neurons with a purine nucleoside, or analog thereof. Preferably, inosine or guanosine is used to stimulate axonal outgrowth and 6-thioguanine is used to inhibit axonal outgrowth. The methods and compns. are particularly useful for modulating the axonal outgrowth of mammalian central nervous system neurons, such as mammalian retinal ganglion cells. Pharmaceutical and packaged formulations that include the purine nucleosides, and analogs thereof, of the invention are also provided. 56-65-5, 5'-ATP, biological studies 58-64-0, 5'-ADP, ΤT biological studies RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (purine nucleosides and analogs for modulating the axonal outgrowth of central nervous system neurons) RETABLE Referenced Author | Year | VOL | PG | Referenced Work | Referenced |(RPY)|(RVL)|(RPG)| (RWK) | File (RAU) \_ |J NEUROSCI Greene, L |1990 |10 | HCAPLUS | INT J DEV NEUROSCI | HCAPLUS Gysbers, J |1996 |14 1997 | NEUROREPORT | HCAPLUS Gysbers, J |1992 |3 IWO 9400132 A Medcament, P |1994 | HCAPLUS 

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|EUR J NEUROSCI
Svensson, B
                      |1993 |5
                                                           MEDLINE
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L130 ANSWER 21 OF 31 HCAPLUS COPYRIGHT 2002 ACS
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1999:166635 HCAPLUS

DN

- ΤI Compositions, kits, and methods for effecting adenine nucleotide modulation of DNA mismatch recognition proteins
- ΙN Fishel, Richard; Gradia, Scott; Acharya, Samir
- PΑ Thomas Jefferson University, USA
- SO PCT Int. Appl., 165 pp.

CODEN: PIXXD2

DТ Patent

LA English

FAN.	CNT	1																
	PAT	ENT :	NO.		KI	ND	DATE			APPLICATION NO. DATE					,			
														1000000				
ΡI	WO	9910			<b>A</b>		1999					98-U						
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			DK,	EE,	ES,	FI,	GB,	GE,	GH,	GM,	HR,	ΗU,	ID,	IL,	IS,	JP,	ΚE,	KG,
			ŔΡ,	KR,	KZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD,	MG,	MK,	MN,	MW,	MX,
			NO,	NZ,	PL,	PΤ,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	TT,
			UA,	ŪG,	US,	UZ,	VN,	YU,	ZW,	AM,	AZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM
		RW:	GH,	GM,	KE,	LS,	MW,	SD,	SZ,	UG,	ZW,	AT,	BE,	CH,	CY,	DE,	DK,	ES,
			FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,	CI,
			CM,	GA,	GN,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG						
	ΑU	9891	251		Α	1	1999		AU 1998-91251 19980828									
	US	6333	153		В	1	2001	1225		US 1998-143571					19980828			
	US	2002	0582	75	Α	1	2002	0516		U:	S 20	01-9	3490	9	2001	0822		
PRAI	US	1997	-571	36P	Р		1997	0828										
	US	1997	-669	77P	P		1997	1128										
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	US US US	JS 1997-66977P JS 1998-93935P JS 1998-143571		P P A		1997 1998 1998 1998	1128 0723 0828											

AΒ Compns., and products comprising a MutS homolog which binds to a mismatched region of a duplex DNA mol. in the presence of ADP are provided, as are methods of binding MutS homologs to mismatched DNA in the presence of ADP. Heterodimers of hMSH2:hMH6 are demonstrated to act as a mol. switch which is activatable by ADP based on the activity in binding to mismatched duplex DNA, ATPase activity, and function in mismatch The use of MutL homolog derivs. in combination with MutS homologs is also included. Mutations obsd. in hMSH2 affect the interaction of hMSH2 with other MutS homologs and are assocd. with hereditary non-polyposis colon cancer. Purified human MSH5 homolog and its cDNA sequence are also provided. Finally, nonhuman mammals which are nullizygous for both Msh2 and p53 are also provided, as are methods of making and using the same. The compns. of the present invention have applications as diagnostic reagents in deg. whether a compd. affects tumorigenesis, apoptosis, aging, fetal development, and gene expression of p53 and MutS.

IT 56-65-5, 5'-ATP, biological studies 58-64-0, 5'-ADP,

biological studies

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(adenine nucleotide modulation of DNA mismatch recognition protein MutS and MutL homologs)

#### RETABLE

Referenced Author (RAU)			(RPG)		Referenced   File
Acharya	•	•	•	Proc Natl Acad Sci	
Chi	11994	1269	29984	J Biol Chem	HCAPLUS
Chi	11994	1269	129993	J Biol Chem	HCAPLUS
Drummond	11995	1268	1909	Science	HCAPLUS
Fearon	11990	161	1759	Cell	HCAPLUS
Fishel	11994	184	5539	Cancer Res	
Fishel	11993	175	1027	Cell	HCAPLUS
Fishel	11994	1266	11403	Science	HCAPLUS

L130 ANSWER 22 OF 31 HCAPLUS COPYRIGHT 2002 ACS

AN 1998:744969 HCAPLUS

DN 130:20593

TI The use of biologically active substances for influencing the extracellular space of sensory cells

IN Eckmiller, Marion Sangster

PA Germany

SO PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DT Patent

LA German

FAN.CNT 1

- TAIV.	PATENT NO.					ND	DATE			A	PPLI	CATI	и ис	٥.	DATE			
PI	PI WO 9850065 WO 9850065			A A		1998			W	0 19	98-E	P195	1	1998	0402			
	WO	W:	AM,	-	AU,	BA,	BB,	BG,							CZ,			
			LR,	LT,	LU,	LV,	MD,	MG,	MN,	MW,	MX,	NO,	NZ,	PL,	KR, PT,	RO,	RU,	SD,
			•	•	•	•	SL, TJ,	•	TT,	UA,	US,	UZ,	VN,	YU,	ZW,	AM,	AZ,	BY,
		RW:													CY, BJ,			
			CM,	GA,	GN,	ML,	MR,	NE,	SN,	TD,	TG						•	
		1971				_	1998				-	97-1			19970505			
	ΑU	9876	417		Α	1	1998	1127		A	U 19	98-7	6417		1998			
	EΡ	980256		Α	2	2000	0223		E	P 19	98-9	2409	7	1998	0402			

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R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
PRAI DE 1997-19718826
                         19970505
     WO 1998-EP1951
                            19980402
AΒ
     The invention relates to the use of an active substance influencing the
     calcium homeostasis of cells to treat degeneration of sensory cells and
     adjacent cells. The effect of higher Ca concns. with and without calpain
     inhibitors on the structure of retinal outer segments was detd.
     56-65-5, ATP, biological studies 58-64-0, ADP,
TΤ
     biological studies
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (drugs for influencing extracellular area of sensory cells)
L130 ANSWER 23 OF 31 HCAPLUS COPYRIGHT 2002 ACS
AN
     1998:550430 HCAPLUS
DN
     129:175919
     Preparation of dinucleotides and their use as modulators of mucociliary
TΙ
     clearance and ciliary beat frequency
IN
     Pendergast, William; Yerxa, Benjamin R.; Rideout, Janet L.; Siddiqi,
     Suhaib M.
     Inspire Pharmaceuticals, Inc., USA
PΑ
SO
     PCT Int. Appl., 49 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 4
                                                            DATE
     PATENT NO.
                                           APPLICATION NO.
                     KIND DATE
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                                                           ______
                                           WO 1998-US2702
                                                            19980206
     WO 9834942
                      A2
                            19980813
PI
     WO 9834942
                      A3
                            20000106
            AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG,
             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
             NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
             UA, UG, US, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
             FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM,
             GA, GN, ML, MR, NE, SN, TD, TG
                            19990504
     US 5900407
                      Α
                                           US 1997-797472
                                                            19970206
                                           US 1997-798508
                                                            19970210
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                       Α
                            19981117
                                           AU 1998-63242
                                                            19980206
     AU 9863242
                       A1 ·
                            19980826
     AU 738907
                       B2
                            20010927
     EP 981534
                       A2
                            20000301
                                           EP 1998-907435
                                                            19980206
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
         R:
             IE, FI
                                           BR 1998-7169
                                                            19980206
     BR 9807169
                            20000606
                       Α
     JP 2001526635
                       T2 "
                            20011218
                                           JP-1998-535055
                                                            19980206
                            20020219
                                           US 1998-101395
                                                            19980710
     US 6348589
                       В1
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                       Α
                            19991006
                                           NO 1999-3776
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     US 2002082417
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                                           US 2001-7451
                                                            20011106
                       Α1
PRAI US 1997-797472
                       A2
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     US 1997-798508
                       A2
                            19970210
     WO 1998-US2702
                       W
                            19980206
                            19980710
     US 1998-101395
                       Α1
     MARPAT 129:175919
OS
GI
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The present invention relates to certain novel dinucleotides I (X = O, AB CH2, imido, CF2; B, B1 = independently nucleobase; Z, Z1 = independently OH, N3; Y, Y1 = independently H, OH; Q = (HPO3)m; n = 0-2; m = 0-2; n + m= 0-4) and formulations thereof which are highly selective agonists of the P2Y2 and/or P2Y4 purinergic receptor. They are useful in the treatment of chronic obstructive pulmonary diseases such as chronic bronchitis, PCD, cystic fibrosis, as well as prevention of pneumonia due to immobility. Furthermore, because of their general ability to clear retained mucus secretions and stimulate ciliary beat frequency, the compds. of the present invention are also useful in the treatment of sinusitis, otitis media and nasolacrimal duct obstruction. They are also useful for treatment of dry eye disease and retinal detachment. Thus, P1, P2-di(uridine-5'-)-P2, P3-methylenetetraphosphate was prepd. as P2Y2 and/or P2Y4 purinergic receptor (EC50 = 11.1 .mu.mol). 2596-55-6P 4130-19-2P 5542-28-9P IT 5959-90-0P 6674-45-9P 10527-46-5P 10527-48-7P 13457-68-6P 26184-65-6P 30632-08-7P 34692-44-9P 41708-91-2P 56983-23-4P 59985-20-5P 59985-21-6P 61340-12-3P 63785-59-1P 79695-24-2P 79695-25-3P 83008-69-9P 88109-92-6P 96920-51-3P 97776-54-0P 97776-55-1P 103137-88-8P 103137-89-9P 111035-55-3P 111648-11-4P 134311-47-0P 135780-83-5P 135780-85-7P 135780-92-6P 135802-64-1P 154960-70-0P 170638-56-9P 170638-57-0P 170638-58-1P 170638-59-2P 170638-60-5P 170638-61-6P 170638-62-7P 211427-06-4P 211427-09-7P 211427-11-1P 211448-67-8P 211448-72-5P 211448-73-6P 211448-74-7P 211448-76-9P 211448-77-0P 211448-78-1P 211448-79-2P 211448-80-5P 211448-81-6P

211448-88-3P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of dinucleotides and their use as modulators of mucociliary clearance and ciliary beat frequency)

TT 63-39-8, Uridine 5'-triphosphate 19817-92-6 211448-71-4

RL: RCT (Reactant); RACT (Reactant or reagent)

(prepn. of dinucleotides and their use as modulators of mucociliary clearance and ciliary beat frequency)

IT 211427-07-5P 211448-70-3P 211448-75-8P 211448-82-7P 211448-83-8P 211448-84-9P 211448-85-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of dinucleotides and their use as modulators of mucociliary clearance and ciliary beat frequency)  $\,$ 

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AN
     1998:548515 HCAPLUS
     129:180137
DN
TI
     Method of treating dry eye disease with purinergic receptor agonists
TN
     Yerxa, Benjamin R.; Jacobus, Karla M.; Pendergast, William; Rideout, Janet
PΑ
     Inspire Pharmaceuticals, Inc., USA
SO
     PCT Int. Appl., 41 pp.
     CODEN: PIXXD2
DΤ
     Patent
LA
     English
FAN.CNT 4
                      KIND DATE
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     PATENT NO.
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                            19980813
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                      Α1
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             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
             NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
             UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
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             GA, GN, ML, MR, NE, SN, TD, TG
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                       B2
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     ZA 9800979
                       Α
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     EP 1003474
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                                                            19990804
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                            20020530
                                           US 2001-10055
                                                            20011109
     US 2002065244
PRAI US 1997-797472
                       A2
                            19970206
                       W
                            19980206
     WO 1998-US2701
                            19981.014
     US 1998-171169
                       Α1
     MARPAT 129:180137
OS
     A method and prepn. for the stimulation of tear secretion in a subject in
AΒ
     need of such treatment is disclosed. The method comprises administering
     to the ocular surfaces of the subject a purinergic receptor agonist such
     as UTP, dinucleotides, CTP, ATP, or their therapeutically useful analogs
     and derivs., in an amt. effective to stimulate tear fluid secretion and
     enhance drainage of the lacrimal system. Pharmaceutical formulations and
     methods of making the same are also disclosed. Methods of administering
     the same would include: topical administration via a liq., gel, cream, or
     as part of a contact lens or selective release membrane; or systemic
     administration via nasal drops or spray, inhalation by nebulizer or other
     device, oral form (liq. or pill), injectable, intra-operative instillation
     or suppository form. P1,P4-di(uridine tetraphosphate) tetrasodium salt
     (I) was formulated as an isotonic aq. soln. and topically administered to
     the eyes of rabbits; I at 0.5, 5.0, and 8.5 % concn. significantly
     increased tear secretion.
     56-65-5, ATP, biological studies 63-39-8, UTP
IT
     65-47-4, CTP 211427-06-4 211427-07-5
     211427-08-6 211427-09-7 211427-10-0
     211427-11-1
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (dry eye disease treatment with purinergic receptor agonists)
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L130 ANSWER 25 OF 31 HCAPLUS COPYRIGHT 2002 ACS

1998:248716 HCAPLUS

ΑN

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DN 128:248610
TI Composition
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TI Compositions containing cereb for treating optic atrophy

IN Hu, Chang; Yu, Dongsheng; Wang, Lin

PA Gete Biological Engineering Science & Technology Development Co., Beijing, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 7 pp. CODEN: CNXXEV

DT Patent

LA Chinese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	CN 1138985	A	19970101	CN 1995-104482	19950628
	CN 1036119	В	19971015		

AB Compns. [freeze-dried injections] for treating optic atrophy contain cereb and inosine, ATP and/or CoA at 3-7: 0.5-2: 0.1-3: 0.1-2.5. Effectiveness was clin. tested.

IT 56-65-5, ATP, biological studies

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (compns. contg. cereb for treating optic atrophy)

L130 ANSWER 26 OF 31 HCAPLUS COPYRIGHT 2002 ACS

AN 1998:152041 HCAPLUS

DN 128:261811

TI Preparation of liposome-encapsulating adenosine triphosphate

AU Arakawa, Akira; Ishiguro, Sei-Ichi; Ohki, Kazuo; Tamai, Makoto

CS Department Ophthalmology, Tohoku University School Medicine, Sendai, 980-8574, Japan

SO Tohoku Journal of Experimental Medicine (1998), 184(1), 39-47 CODEN: TJEMAO; ISSN: 0040-8727

PB Tohoku University Medical Press

DT Journal

LA English

AB Liposomes encapsulating ATP (ATP) were prepd. by sonication, and the liposomes were evaluated for use in a drug delivery system. The liposomes, which were composed of phosphatidylcholine and cholesterol, were about 1.1 .mu.m in size, as obsd. under a microscope. From their size, the vesicles were thought to be multilamellar. The max. concn. of ATP in the liposomes was 1.0 mM, when the initial concns. of lipid and ATP were 20 mM and 300 mM, resp. The max. entrapment ratio of ATP in the liposomes was 88%, when the initial concns. of lipid and ATP were 20 mM and 500 mM, resp. About 4% of ATP was encapsulated in these expts. When liposomes contained 4-7% of cholesterol, about 35% of encapsulated ATP was released from the liposomes for 90 h at 37.degree.C in vitro. These findings indicated that liposomes encapsulating ATP could be used for the treatment of ischemic retina.

IT 56-65-5, Atp, biological studies
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PEP (Physical, engineering or chemical process);
THU (Therapeutic use); BIOL (Biological study); PROC (Process);
USES (Uses)

(prepn. of liposome-encapsulated ATP)

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L130 ANSWER 27 OF 31 HCAPLUS COPYRIGHT 2002 ACS
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AN 1998:38360 HCAPLUS

DN 128:93211

TI Medicament base for local use in the eyes

PA Schrage, Norbert, Germany

SO Ger. Offen., 4 pp. CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

- PI DE 19626479 A1 19980108 DE 1996-19626479 19960702
- AB A phosphate-free buffered isotonic soln. for application to the cornea is provided for buffering and neutralizing corrosive or irritating substances in the eye. The soln. contains the same proportions of electrolytes as the cornea as well as sugars, polyols, polysaccharides, amino acids, and org. buffers such as maleate, citrate, acetate, HEPES, or Tris. Phosphates are omitted to prevent calcification of the cornea.
- L130 ANSWER 28 OF 31 HCAPLUS COPYRIGHT 2002 ACS
- AN 1997:472375 HCAPLUS
- DN 127:130966
- TI Effects of adrenergic agents and phosphodiesterase inhibitors on outflow facility and cell shape of bovine trabecular meshwork
- AU Suzuki, Ryo; Karageuzian, Levon N.; Crean, Edmund V.; Anderson, P. John
- CS Department of Ophthalmology, Yamaguchi University School of Medicine, Ube, 755, Japan
- SO Japanese Journal of Ophthalmology (1997), 41(1), 31-37 CODEN: JJOPA7; ISSN: 0021-5155
- PB Elsevier
- DT Journal
- LA English
- Changes in the outflow facility of perfused calf eyes and in the shape of AΒ cells in cultured trabecular meshwork (TM) have been studied, following exposure to adrenergic agents and phosphodiesterase inhibitors (PDE). Dobutamine caused confluent TM cells to change their usual polygonal shape to a characteristic stellate shape. Salbutamol had no effect, but PDE inhibitors, isobutylmethylxanthine (IBMX), theophylline, and caffeine were very effective in producing this shape change. Epinephrine, isoproterenol, dobutamine, and salbutamol did not increase the outflow facility, either at 22.degree. or 36.degree., while theophylline, caffeine, and IBMX did increase it in a dose-dependent manner. The high concns. of .beta.-adrenergic agents required to produce even a small change in outflow facility and cell shape argue against the involvement of adrenergic-receptor mediation and may suggest another mechanism; on the other, the enhancement of epinephrine effects by PDE inhibitors and the similar effect produced by cyclic adenosine 3',5'-cyclic phosphate (cAMP) and purines suggest that changes in the cell shape are produced by .beta.-receptor activation. The .beta.-adrenergic agents were not effective in changing outflow facility, but the PDE inhibitors were remarkably effective both in changing the shape and in increasing facility. The results are discussed in relation to treatment of glaucoma.
- IT 56-65-5, 5'-ATP, biological studies 58-64-0, 5'-ADP, biological studies
  - RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(effects of adrenergic agonists and phosphodiesterase inhibitors on outflow facility and cell shape of bovine trabecular meshwork in relation to **glaucoma** treatment)

- L130 ANSWER 29 OF 31 HCAPLUS COPYRIGHT 2002 ACS
- AN 1997:209102 HCAPLUS
- DN 126:275487
- TI Extracellular ATP activates calcium signaling, ion, and fluid transport in retinal pigment epithelium
- AU Peterson, Ward M.; Meggyesy, Chris; Yu, Kefu; Miller, Sheldon S.
- CS School of Optometry and Department of Molecular and Cell Biology,

```
University of California, Berkeley, CA, 94720, USA Journal of Neuroscience (1997), 17(7), 2324-2337 CODEN: JNRSDS; ISSN: 0270-6474
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- PB Society for Neuroscience
- DT Journal
- LA English
- AB The presence of receptors for ATP has not been established in any native prepn. of retinal neurons or glia. In the present study, we used conventional electrophysiol. and [Ca2+]in fluorescence imaging techniques to investigate the effects of ATP added to Ringer's soln. perfusing the retinal-facing (apical) membrane of freshly isolated monolayers of bovine retinal pigment epithelium (RPE). ATP (or UTP) produced large, biphasic voltage and resistance changes with a Kd of .apprx.5 .mu.M for ATP and .apprx.1 .mu.M for UTP. Elec. and pharmacol. evidence indicates that the first and second phases of the response are attributable to an increase in basolateral membrane Cl conductance and a decrease in apical membrane K conductance, resp. The ATP-induced responses were not affected by adenosine, but were reduced by the P2-purinoceptor blocker suramin. ATP also produced a large, transient increase in [Ca2+]in that was blocked by cyclopiazonic acid, an inhibitor of endoplasmic reticulum Ca2+-ATPases. The calcium buffer BAPTA attenuated the voltage effects of ATP. We also found that apical DIDS significantly inhibited the ATP-evoked [Ca2+]in and elec. responses, suggesting that DIDS blocked the purinoceptor. Measurements of fluid movement across the RPE using the capacitance probe technique demonstrated a significant increase in fluid absorption by apical UTP. These data indicate the presence of metabotropic P2Y/P2U-purinoceptors at the RPE apical membrane and implicate extracellular ATP in vivo as a retinal signaling mol. that could help regulate the hydration and chem. compn. of the subretinal space.
- IT 56-65-5, 5'-ATP, biological studies
  RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)
  (extracellular ATP activates calcium signaling, ion, and fluid transport in retinal pigment epithelium)
- IT 63-39-8, UTP

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(extracellular ATP activates calcium signaling, ion, and fluid transport in retinal pigment epithelium in relation to)

L130 ANSWER 30 OF 31 HCAPLUS COPYRIGHT 2002 ACS

AN 1996:537788 HCAPLUS

DN 125:158638

TI Method of treatment of diabetes mellitus by administration of adenosine 5'-triphosphate and other adenine nucleotides

IN Rapaport, Eliezer

PA USA

SO U.S., 6 pp. CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI US 5547942 A 19960820 US 1994-17771 19940104

AB The administration of adenine nucleotides or adenosine and inorg. phosphate to a human patient results in the generation of elevated liver, other organ, and red blood cell ATP pools as well as increased levels of ATP in the extracellular blood plasma compartment of the blood. The present invention deals with the utilization of the elevated extracellular levels of ATP for achieving the well-established stimulation of insulin secretion following the interactions of extracellular ATP pools with pancreatic .beta. cell purine receptors. The invention is therefore

concerned with the treatment of patients suffering from non-insulin-dependent diabetes mellitus (NIDDM or Type-II diabetes) and their chronic clin. complications which are the result of continuous hyperglycemia, by the administration of these physiol. agents.

IT 56-65-5, Adenosine 5'-triphosphate, biological studies
58-64-0, Adenosine 5'-diphosphate, biological studies
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(ATP and other adenine nucleotides for diabetes treatment)

L130 ANSWER 31 OF 31 HCAPLUS COPYRIGHT 2002 ACS

AN 1995:416436 HCAPLUS

DN 122:170250

TI Pharmaceutical compositions containing inhibitors of proteic ADP ribosylation are useful to prevent the diabetes mellitus complications

IN Gorio, Alfredo; Borella, Fabio

PA Instituto Biochimico Italiano Giovanni Lorenzini S.p.A., Italy

SO Eur. Pat. Appl., 17 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI EP 638309 A1 19950215 EP 1994-110805 19940712 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE

PRAI IT 1993-MI1554 19930714

AB Pharmaceutical compns. contg. inhibitors of proteic ADP ribosylation are useful to prevent the diabetes mellitus complications such as neuropathies, nephropathies, retinopathies, macroangiopathies, microangiopathies, and hepatopathies. The effectiveness of vitamin K1 in decreasing blood glucose level of diabetic rats is reported. A hard gelatin pearl contained vitamin K1 10, lactose 62, maize starch 27, and Mg stearate lmg.

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TSCA INFORMATION NOW CURRENT THROUGH January 7, 2002

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

VAR G1=O/S
VAR G2=O/CH2/NH/11
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 12

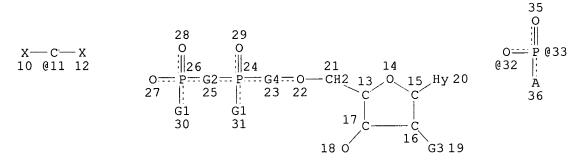
STEREO ATTRIBUTES: NONE

L12 24293 SEA

24293 SEA FILE=REGISTRY SSS FUL L10

L13

STR



VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
REP G4=(0-1) 32-24 33-22
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 20
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

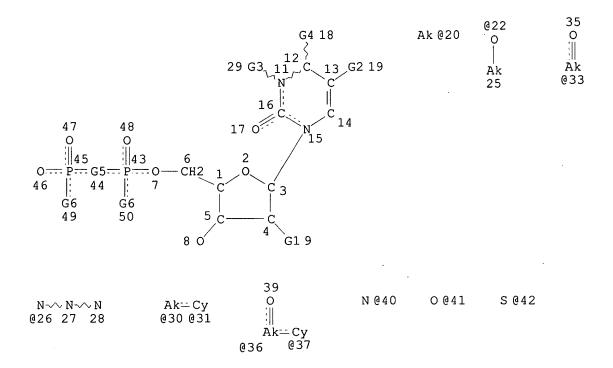
GRAPH ATTRIBUTES: RING(S) 'ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 26

STEREO ATTRIBUTES: NONE

L15 4525 SEA FILE=REGISTRY SUB=L12 CSS FUL L13

L18 STR

Jan Delaval Reference Librarian Biotechnology & Chemical Library CM1 1E07 – 703-308-4498 jan.delaval@uspto.gov



X-C-X 51 @52 53

VAR G1=H/O VAR G2=H/X/AK/20/22/NO2/26 VAR G3=H/AK/30/31/33/36/37 VAR G4=41/42/40VAR G5=O/N/CH2/52 VAR G6=O/S NODE ATTRIBUTES: CONNECT IS M1 RC AT 20 40 CONNECT IS M1 RC AT CONNECT IS M1 RC AT 41 CONNECT IS M1 RC AT 42 CONNECT IS M1 RC AT 43 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 46

STEREO ATTRIBUTES: NONE

L20 246 SEA FILE=REGISTRY SUB=L15 CSS FUL L18

L24 ST

N--- G2 G2--- N--- G2 660 61 62 663 64

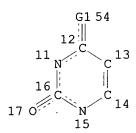
VAR G1=0/S/N/55/57/60/63 VAR G2=AK/20/CY/59/30 NODE ATTRIBUTES: CONNECT IS M1 RC AT 11 CONNECT IS M1 RC AT 13 15 CONNECT IS M1 RC AT 20 CONNECT IS M1 RC AT CONNECT IS M1 RC AT 59 DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 21

STEREO ATTRIBUTES: NONE

L26 240 SEA FILE=REGISTRY SUB=L20 CSS FUL L24 L27 STR



VAR G1=O/S
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 11
CONNECT IS M1 RC AT 13
CONNECT IS M1 RC AT 15
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 8

STEREO ATTRIBUTES: NONE

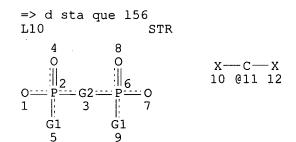
L28 156 SEA FILE=REGISTRY SUB=L26 CSS FUL L27

L29 84 SEA FILE=REGISTRY ABB=ON PLU=ON L26 NOT L28

L30 224 SEA FILE=REGISTRY ABB=ON PLU=ON (L26 OR L28 OR L29) NOT (11C# OR 13C# OR 14C# OR C11# OR C13# OR C14# OR (D OR T)/ELS

OR 32P)

L31	211	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L30	NOT	IDS/CI
L32	192	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L31	NOT	COMPD
L33	184	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L32	NOT	PMS/CI
L34	182	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L33	NOT	33S
L35	180	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L34	NOT	36S
L36	178	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L35	NOT	32S
L37	166	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L36	NOT	(LABELED OR 32P2 OR
		180‡	OR 34S)					
L40	165	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L37	NOT	FERROCENYL



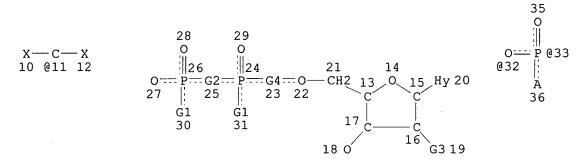
VAR G1=O/S
VAR G2=O/CH2/NH/11
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

L12 24293 SEA FILE=REGISTRY SSS FUL L10

L13 STR



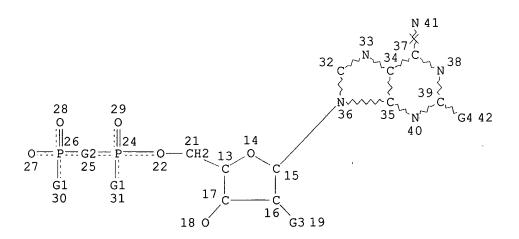
VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
REP G4=(0-1) 32-24 33-22
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 20
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 26

STEREO ATTRIBUTES: NONE

L15 4525 SEA FILE=REGISTRY SUB=L12 CSS FUL L13



VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
VAR G4=H/X/43
NODE ATTRIBUTES:
NSPEC IS RC AT 41
CONNECT IS M1 RC AT 38
CONNECT IS M1 RC AT 41
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 33

STEREO ATTRIBUTES: NONE

L41

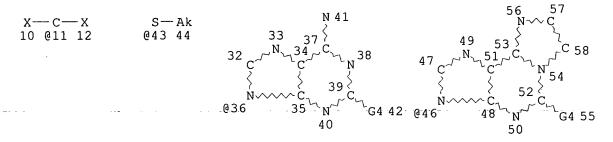
SCR 2039 OR 2043 OR 2048

L43

334 SEA FILE=REGISTRY SUB=L15 CSS FUL L38 NOT L41

L44

STR



```
VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
VAR G4=H/X/43
VAR G5=36/46/59
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED
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### GRAPH ATTRIBUTES:

RSPEC I

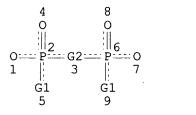
NUMBER OF NODES IS 59

### STEREO ATTRIBUTES: NONE

L46	266	SEA	FILE=REGISTRY	SUB=L43	CSS FUL	L44	
L47	190	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L46 NOT	COMPD
L48	8	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L47 AND	NC>=2 NOT SALT
L50	137	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L47 AND	NC>=2 NOT L48
L51	132	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L50 NOT	MXS/CI
L52	12	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L51 AND	(C28H22N2 OR
		C21	H20N3 OR C15H1	6N3 OR C	15H14N OI	R C15H11	N2 OR C36H24N4 OR
		C44	H38N8 OR C122H	200N12 OI	R C46H681	N4 OR C3	8H30N4 OR C21H15N2)
L53	5	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L51 AND	CU/ELS
L54	3	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L53 NOT	KAPPA
L55	2	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L53 NOT	L54
L56	118	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L51 NOT	(L52 OR L55)

=> =>

=> d sta que 192 L10 STR



VAR G1=O/S
VAR G2=O/CH2/NH/11
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 12

# STEREO ATTRIBUTES: NONE

L12 24293 SEA FILE=REGISTRY SSS FUL L10 L41 SCR 2039 OR 2043 OR 2048

L88 STR

068

Ak @79

 $N \sim N \sim N$  @76 77 78

VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
VAR G4=36/45/57/68
VAR G5=AK/74/NO2/X/76/79
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 79
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED

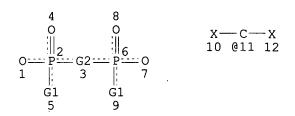
RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 67

STEREO ATTRIBUTES: NONE

L91 38 SEA FILE=REGISTRY SUB=L12 CSS FUL L88 NOT L41 L92 34 SEA FILE=REGISTRY ABB=ON PLU=ON L91 NOT COMPD

=> =>

=> d sta que 176 L10 STR



VAR G1=O/S VAR G2=O/CH2/NH/11 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

L12 24293 SEA FILE=REGISTRY SSS FUL L10

L13 STR

35 0 28 29 0 0 O--- P @33 26 24 @32 10 @11 12 21 O-P-G2-P-G4-O--CH2 13 Hy 20 Ä 25 23 22 27 36 G1 G1 17 30 31 16 G3 19 18 O

VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
REP G4=(0-1) 32-24 33-22
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 20
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 26

STEREO ATTRIBUTES:- NONE

L15 4525 SEA FILE=REGISTRY SUB=L12 CSS FUL L13

L24 STR

N--- G2 G2--- N--- G2 660 61 62 063 64

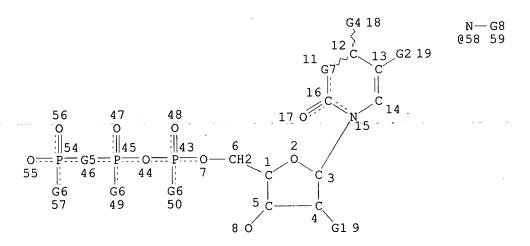
VAR G1=0/S/N/55/57/60/63 VAR G2=AK/20/CY/59/30 NODE ATTRIBUTES: CONNECT IS M1 RC AT 13 CONNECT IS M1 RC AT CONNECT IS M1 RC AT 15 CONNECT IS M1 RC AT 20 CONNECT IS M1 RC AT DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 21

STEREO ATTRIBUTES: NONE

L41 SCR 2039 OR 2043 OR 2048

L64 STR



VAR G1=H/O VAR G2=H/X VAR G4=41/42/40 VAR G5=O/N/CH2/52

```
VAR G6=O/S
VAR G7=N/58
VAR G8=AK/33/20/30/31
NODE ATTRIBUTES:
CONNECT IS M1 RC AT
                      20
CONNECT IS M1
               RC AT
                      40
CONNECT IS M1
               RC AT
                      41
CONNECT IS M1
              RC AT
                      42
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED
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## GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

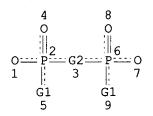
NUMBER OF NODES IS 43

### STEREO ATTRIBUTES: NONE

L66	271	SEA	FILE=REGISTRY	SUB=L15	CSS FUL	L64 NOT	L41
L68	257	SEA	FILE=REGISTRY	SUB=L66	CSS FUL	L24	
L69	14	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L66 NOT	. r
L70	2	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L69 AND	(C10H18N3O14P3 OR
		C101	118N3O15P3)				
L71	255	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L68 NOT	MXS OR IDS)/CI
L72	230	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L71 NOT	COMPD
L73	81	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L72 AND	NC>=2
L74	77	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L73 NO	CCS/CI
L75	· 149	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L72 NOT	L73
L76	228	SEA	FILE=REGISTRY	ABB=ON	PLU=ON	(L70 OF	R L74 OR L75)

=> =>

=> d sta que 187 L10 STR



X-C-X 10 @11 12

VAR G1=O/S
VAR G2=O/CH2/NH/11
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

# GRAPH ATTRIBUTES:

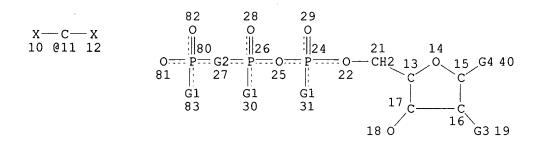
RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 12

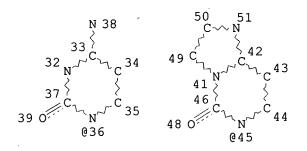
### STEREO ATTRIBUTES: NONE

L12 24293 SEA FILE=REGISTRY SSS FUL L10

L41 SCR 2039 OR 2043 OR 2048

L80 STR





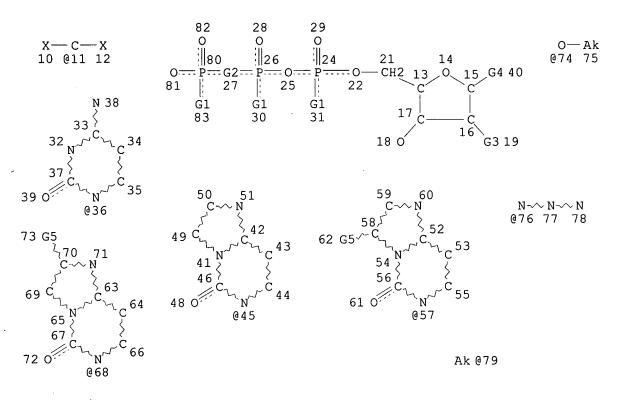
VAR G1=O/S VAR G2=O/CH2/NH/11 VAR G3=O/H VAR G4=36/45 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 43

STEREO ATTRIBUTES: NONE

L82 488 SEA FILE=REGISTRY SUB=L12 SSS FUL L80

L83 STR



VAR G1=O/S
VAR G2=O/CH2/NH/11
VAR G3=O/H
VAR G4=36/45/57/68
VAR G5=AK/74/NO2/S/76/79
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 79
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 71

STEREO ATTRIBUTES: NONE

=>

L86 61 SEA FILE=REGISTRY SUB=L82 CSS FUL L83 NOT L41 L87 50 SEA FILE=REGISTRY ABB=ON PLU=ON L86 NOT COMPD

VAR G1=O/S VAR G2=O/CH2/NH/11 NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

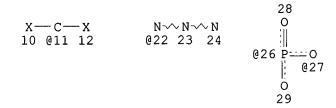
NUMBER OF NODES IS 12

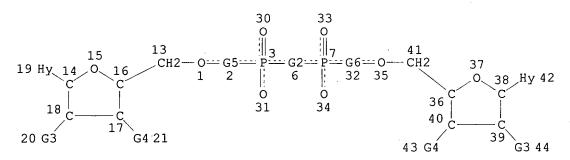
STEREO ATTRIBUTES: NONE

L12 24293 SEA FILE=REGISTRY SSS FUL L10

L41 SCR 2039 OR 2043 OR 2048

L93 STR





VAR G2=O/CH2/11/NH

VAR G3=H/O

VAR G4=0/22

REP G5=(0-2) 26-1 27-3

REP G6=(0-2) 27-7 26-35

NODE ATTRIBUTES:

CONNECT IS M1 RC AT 19

CONNECT IS M1 RC AT 42

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

### GRAPH ATTRIBUTES:

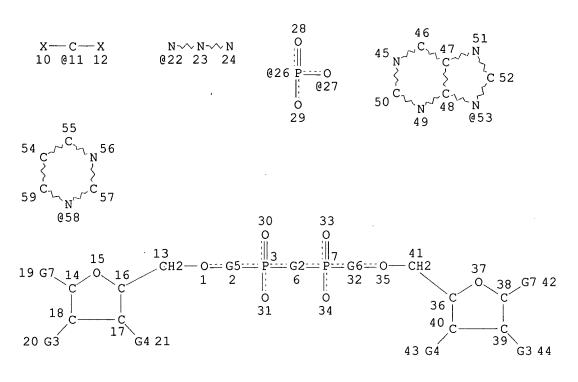
RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 39

STEREO ATTRIBUTES: NONE

L96 1242 SEA FILE=REGISTRY SUB=L12 CSS FUL L93 NOT L41

L97 STR



VAR G2=O/CH2/11/NH

VAR G3=H/O

VAR G4=0/22

REP G5=(0-2) 26-1 27-3

REP G6=(0-2) 27-7 26-35

VAR G7=53/58

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

## GRAPH ATTRIBUTES:

RSPEC 51 54

NUMBER OF NODES IS 54

### STEREO ATTRIBUTES: NONE

L98 33

333 SEA FILE=REGISTRY SUB=L96 SSS FUL L97

L99

L1

333 SEA FILE=REGISTRY ABB=ON PLU=ON L98 NOT (MXS OR IDS OR

CCS)/CI

L100

324 SEA FILE=REGISTRY ABB=ON PLU=ON L99 NOT COMPD

# => d his

(FILE 'HOME' ENTERED AT 09:48:11 ON 16 JUL 2002) SET COST OFF

FILE 'HCAPLUS' ENTERED AT 09:48:51 ON 16 JUL 2002

E PETERSON W/AU

26 S E3,E14,E33,E34

L2 5 S L1 AND (1 OR 63)/SC, SX NOT MORPHINE

L3 4 S L1 AND (2 OR 12 OR 13)/SC, SX

SEL RN L2

FILE 'REGISTRY' ENTERED AT 09:53:25 ON 16 JUL 2002

L4 129 S E1-E129

L5 53 S L4 AND P/ELS

FILE 'HCAPLUS' ENTERED AT 09:56:35 ON 16 JUL 2002 SEL RN L3

```
FILE 'REGISTRY' ENTERED AT 09:56:45 ON 16 JUL 2002
             12 S E130-E141
L7
L8
              3 S L7 AND P/ELS
L9
              2 S L8 AND P>=2
L10
                STR
             50 S L10
L11
          24293 S L10 FUL
L12
                SAV TEMP L12 OWENS570/A
L13
                STR L10
L14
             50 S L13 CSS SAM SUB=L12
           4525 S L13 CSS FUL SUB=L12
L15
                SAV TEMP L15 OWENS570A/A
L16
                STR
             50 S L16 CSS SAM SUB=L15
L17
L18
                STR L16
L19
             11 S L18 CSS SAM SUB=L15
L20
            246 S L18 CSS FUL SUB=L15
                SAV L20 TEMP OWENS570AA/A
                STR L18
L21
L22
              0 S L21 CSS SAM SUB=L20
L23
              0 S L21 CSS FUL SUB=L20
L24
                STR L21
             11 S L24 CSS SAM SUB=L20
L25
            240 S L24 CSS FUL SUB=L20
L26
                SAV L26 TEMP OWENS570AAA/A
L27
                STR L24
L28
            156 S L27 CSS FUL SUB=L26
                SAV L28 OWENS570AAAA/A TEMP
             84 S L26 NOT L28
L29
            224 S L26, L28, L29 NOT (11C# OR 13C# OR 14C# OR C11# OR C13# OR C14#
            211 S L30 NOT IDS/CI
L31
L32
            192 S L31 NOT COMPD
L33
            184 S L32 NOT PMS/CI
L34
            182 S L33 NOT 33S
            180 S L34 NOT 36S
L35
            178 S L35 NOT 32S
L36
            166 S L36 NOT (LABELED OR 32P2 OR 180# OR 34S)
                SAV L37 OWEN570AAAAA/A TEMP
L38
                STR L13
             22 S L38 CSS SAM SUB=L15
L39
            165 S L37 NOT FERROCENYL
L40
                SCR 2039 OR 2043 OR 2048
L41
            18 S L38 NOT L41 CSS SAM SUB=L15
L42
            334 S L38 NOT L41 CSS FUL SUB=L15
L43
                SAV TEMP L43 OWENS570B/A
L44
                STR L38
L45
             17 S L44 CSS SAM SUB=L43
            266 S L44 CSS FUL SUB=L43
L46
                SAV L46 OWENS570BB/A
                DEL OWENS570BB/A
                SAV L46 OWENS570BB/A TEMP
            190 S L46 NOT COMPD
L47
              8 S L47 AND NC>=2 NOT SALT
L48
              3 S L48 AND (H2O OR C4H12N)
            137 S L47 AND NC>=2 NOT L48
            132 S L50 NOT MXS/CI
            12 S L51 AND (C28H22N2 OR C21H2ON3 OR C15H16N3 OR C15H14N OR C15H1
L52
L53
              5 S L51 AND CU/ELS
```

```
L54
              3 S L53 NOT KAPPA
L55
              2 S L53 NOT L54
L56
            118 S L51 NOT L52, L55
                SAV L56 TEMP OWENS570BBB/A
                STR L13
L57
              0 S L57 NOT L41 CSS SAM SUB=L15
L58
              0 S L57 NOT L41 SAM SUB=L15
L59
              0 S L57 CSS SAM SUB=L12
L60
              0 S L57 SAM SUB=L12
L61
             23 S L57 FUL SUB=L12
L62
                SAV L62 TEMP OWENS570C/A
L63
              0 S L57 CSS FUL SUB=L62
                SAV L63 TEMP OWENS570CC/A
L64
                STR L18
L65
             16 S L64 NOT L41 CSS SAM SUB=L15
L66
            271 S L64 NOT L41 CSS FUL SUB=L15
                SAV TEMP L66 OWENS570D/A
             16 S L24 CSS SAM SUB=L66
L67
            257 S L24 CSS FUL SUB=L66
                SAV TEMP L68 OWENS570DD/A
L69
             14 S L66 NOT L68
L70
              2 S L69 AND (C10H18N3O14P3 OR C10H18N3O15P3)
L71
            255 S L68 NOT (MXS OR IDS)/CI
            230 S L71 NOT COMPD
L72
             81 S L72 AND NC>=2
L73
             77 S L73 NOT CCS/CI
L74
            149 S L72 NOT L73
L75
L76
            228 S L70, L74, L75
                SAV TEMP L76 OWENS570DDD/A
                STR L57
              0 S L77 CSS SAM SUB=L12
L78
              0 S L77 SAM SUB=L12
L79
                STR L77
L80
L81
             25 S L80 SAM SUB=L12
            488 S L80 FUL SUB=L12
L82
                SAV L82 OWENS570E/A TEMP
L83
                STR L77
              4 S L83 CSS SAM SUB=L82
L84
              2 S L83 NOT L41 CSS SAM SUB=L82
L85
L86
             61 S L83 NOT L41 CSS FUL SUB=L82
                SAV TEMP L86 OWENS570EE/A
L87
             50 S L86 NOT COMPD
                SAV TEMP L87 OWENS570EEE/A
                STR L57
T.88
              2 S L88 CSS SAM SUB=L12
L89
              2 S L88 NOT L41 CSS SAM SUB=L12
L90
             38 S L88 NOT L41 CSS FUL SUB=L12
L91
               SAV TEMP L91 OWENSCCC/A
             34 S L91 NOT COMPD
L92
            STR L10
L93
             50 S L93 CSS SAM SUB=L12
L94
             50 S L93 NOT L41 CSS SAM SUB=L12
L95
L96
           1242 S L93 NOT L41 CSS FUL SUB=L12
                SAV TEMP L96 OWENS570F/A
L97
                STR L93
            333 S L97 FUL SUB=L96
L98
                SAV TEMP L98 OWENS570FF/A
            333 S L98 NOT (MXS OR IDS OR CCS)/CI
L99
            324 S L99 NOT COMPD
L100
             84 S L100 AND NC>=2
L101
            240 S L100 NOT L101
L102
```

SAV TEMP L100 OWENS570FFF/A

Series.

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FILE 'HCAPLUS' ENTERED AT 12:21:12 ON 16 JUL 2002
L103
           9464 S L40 OR L56 OR L92 OR L76 OR L87 OR L100
L104
               5 S L1 AND L103
L105
          69691 S L6 OR L9
L106
              5 S L1 AND L105
L107
               5 S L104, L106
                E RETINA/CT
                E E3+ALL
L108
          13385 S E2
                 E RETINA/CT
                 E E24+ALL
           2229 S E2
L109
                 E RETINA/CT
                E E35+ALL
            583 S E2
L110
                E RETINA/CT
                E E17+ALL
                E RETINA/CT
                E E20+ALL
            303 S E2
L111
                E MACULAR DEGENERAITON/CT
                E E4+ALL
            581 S E2
L112
                E STAGARDT/CT
                E STARGARDT/CT
                E STARGARDT
L113
              97 S E2-E6
                E BEST/CT
                E GLAUCOMA/CT
                E E4+ALL
           2669 S E5, E4+NT
L114
                E RETINITS/CT
                E RETINITIS/CT
                E E4+ALL
            596 S E2
L115
L116
              1 S E1
                E OPTIC NERVE/CT
                E E3+ALL
L117
           1806 S E2
                E EYE/CT
                E E3+ALL
L118
          56834 S E8, E7+NT
                E E35+ALL
L119
          21818 S E3+NT
             82 S L103 AND L108-L119
L120
L121
            690 S L105 AND L108-L119
             46 S L103 AND (RETINA OR RETINAL OR RETINOPATH? OR MACULAR(L) DEGEN
L122
            321 S L105 AND (RETINA OR RETINAL OR RETINOPATH? OR MACULAR(L) DEGEN
L123
              9 S L103 AND (CYSTOID(L)MACULA?(L)EDEM? OR RETINA?(L)DETACH? OR P
L124
             36 S L105-AND (CYSTOID(L) MACULA?(L) EDEM? OR -RETINA?-(L) DETACH? OR P
L125
            739 S L120-L125
L126
             12 S L103(L) THU/RL AND L126
L127
L128
             26 S L105(L) THU/RL AND L126
L129
             28 S L127, L128
L130
              31 S L107, L129
L131
              5 S L1 AND L130
                 SEL HIT RN
     FILE 'REGISTRY' ENTERED AT 12:38:58 ON 16 JUL 2002
L132
             50 S E1-E50
L133
              41 S L132 NOT S/ELS
L134
              38 S L133 NOT THYMIDINE
             13 S L134 AND (C18H27N4O26P5 OR C18H25N4O20P3 OR C18H26N4O22P4 OR
L135
```

L137	1	S	L134	AND	C18H26N4O23P4	
L138	11	S	L135	NOT	(211448-76-9 OF	R 211427-10-0)
T.139	12	S	T.137	T.138	}	

FILE 'REGISTRY' ENTERED AT 13:09:25 ON 16 JUL 2002

FILE 'HCAPLUS' ENTERED AT 13:09:44 ON 16 JUL 2002 SEL HIT RN L130

FILE 'REGISTRY' ENTERED AT 13:10:04 ON 16 JUL 2002

L140	93	S	E51-1	£143		
L141	81	S	L140	NOT	L139	
L142	43	S	L141	NOT	L132	

FILE 'HCAPLUS' ENTERED AT 13:11:29 ON 16 JUL 2002

FILE 'REGISTRY' ENTERED AT 13:12:00 ON 16 JUL 2002